

GenCore version 5.1.6
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OM nucleic - nucleic search, using sw model

Run on: February 9, 2004, 13:03:17 ; Search time 3909 Seconds
(without alignments)
10381.773 Million cell updates/sec

Title: US-09-990-726-222
Perfect score: 992
Sequence: 1 ggcacgagcagaactagg.....aaaaaaaaaaaaaaaaaaaaa 992

Scoring table: IDENTIFY_NUC
Gapop 10.0 , Gapext 1.0

Searched: 2888711 seqs, 2045481386 residues

Total number of hits satisfying chosen parameters: 5777422

Minimum DB seq length: 0
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Post-processing: Minimum Match 0%
Maximum Match 100%
Listing first 150 summaries

Database :

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41: em_hgtgo_other:

Pred. No. is the number of results predicted by chance to have a

score greater than or equal to the score of the result being printed,
and is derived by analysis of the total score distribution.

SUMMARIES

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VERSION AX358896.1 GI:18675349
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SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE 1
AUTHORS Baker, K.P., Deenoyers, L., Gerritsen, M.E., Goddard, A.,
Godowski, P.J., Grimaldi, J.C., Gurney, A.L., Smith, V., Stephan, J.P.,
Watanabe, C.K. and Wood, W.I.
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JOURNAL Patent: WO 0193983-A 149 13-DEC-2001;
Genentech Inc. (US)
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LOCUS AX362389 992 bp DNA linear PAT 15-FEB-2002
DEFINITION Sequence 149 from Patent WO0208288.
ACCESSION AX362389
VERSION AX362389.1 GI:18694652
KEYWORDS Homo sapiens (human)
SOURCE
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ORGANISM	Homo sapiens
REFERENCE	Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS	Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE	1 Baker, K.P., Desnovers, L., Gerritsen, M.E., Goddard, A.,
JOURNAL	Godowski, P.J., Grimaldi, J.C., Gurney, A.L., Smith, V., Stephan, J.P.,
FEATURES	Watanabe, C.K. and Wood, W.I.
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LOCUS
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ACCESSION
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VERSION
JP 2002010789-A/47.
KEYWORDS
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SOURCE
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Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 505)
Edwards,J.B.D.M., Jobert,S. and Giordano,J.E.
EST and encoded human protein
Patent: JP 2002010789-A 47 15-JAN-2002;
GENSET CORP
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PN JP 2002010789-A/47
PD 15-JAN-2002
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PR 05-AUG-1999 US 60/147499
PI JEAN BAPTISTE DUMAS MILNE EDWARDS, SEVELIN JOBERT, JEAN EVE PI
GIORDANO
PC C12N15/09, C12N15/09, C07K16/47, C07K16/18, C12N1/15, C12N1/19, PC
C12N1/21,

PC C12N5/10, C12P21/02, C12P21/08, C12Q1/68, C12N15/00, C12N5/00, PC
C12N15/00
CC Von Heijne matrix
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CC seq LFCIAVLAASSFS/KK
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Best Local Similarity 99.8%; Pred. No. 6.5e-111;
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Db 61 TGGCTGCCACAGCTTCTCAAGGCGGAGGAGAAATTAACCCCTGCTGCTCCATTG 120
Qy 157 CCTACAAAGTCTTGGAAAGTTTCCCAAGGCGGCTGGGTGCTCATTAACCTGCTGTGCAC 216
Db 121 CCTACAAAGTCTTGGAAAGTTTCCCAAGGCGGCTGGGTGCTCATTAACCTGCTGTGCAC 180
Qy 217 CCCAGCCACACCGCCCATCACTTATTCCTCTGTGTGAAACCAAGATGAGGCA 276
Db 181 CCCAGCCACACCGCCCATCACTTATTCCTCTGTGTGAAACCAAGATGAGGCA 240
Qy 277 AGAAGTGTGAGAGCCCAAGCGGCGCTTCAACCTCAAGCTCAGCTCAGTCAAGTCCA 336
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Db 301 GTCCAGACCTGCTCAGCTTCTTGTGCGGGGCTCCCTCCACCTCAGGTGCCATGTGACA 360
Qy 397 GTCCAGGCTACAGATCAGTGGAGCTGGTCCAAAGCCAGTGTCTGAGTGGGGGCA 456
Db 361 GTCCAGGCTACAGATCAGTGGAGCTGGTCCAAAGCCAGTGTCTGAGTGGGGGCA 420
Qy 457 ACTTCACCTGTCCAGGACAGAGGGGCGGCGGCTCCAGCTGAGATGATCTGCCAGCGTCT 516
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Qy 517 CGGCGAGCCACCTATCACCAACAG 541
Db 481 CGGCGAGCCACCTATCACCAACAG 505
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DEFINITION
Sequence 695 from Patent WO02083898.
ACCESSION
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VERSION
AX588820.1 GI:27900608
KEYWORDS
Homo sapiens (human)
SOURCE
Homo sapiens
ORGANISM
Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1
Bejanin,S., Tanaka,H., Dumas Milne Edwards,J.B., Jobert,S. and
Giordano,J.Y.
Full-length human cdnas encoding potentially secreted proteins
Patent: WO 02083898-A 695 24-OCT-2002;
JOURNAL

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ACCESSION G60502				
VERSION G60502.1 GI:6125821				
KEYWORDS STS.				
SOURCE Homo sapiens (human)				
ORGANISM Homo sapiens				
REFERENCE 1 (bases 1 to 414)				
AUTHORS Myers,R.M.				
TITLE Human STSs (1999)				
JOURNAL Unpublished (1999)				
COMMENT				
Contact: Richard M. Myers				
Stanford Human Genome Center (SHGC)				
Stanford University School of Medicine				
Department of Genetics, M-344, Stanford, CA 94305, USA				
Tel: 4157259687				
Fax: 4157259689				
Email: myers@shgc.stanford.edu				
Primer A: GTGGCAAGAGCAGCTAAAC				
Primer B: GGAATGGGAGGTACAGG				
STS size: 150				
PCR Profile:				
Initial incubation: 95 degrees C for 10 minutes				
Denaturation: 94 degrees C for 30 seconds				
Annealing: 60 degrees C for 30 seconds				
Polymerization: 72 degrees C for 23 seconds				
PCR Cycles: 30				
Thermal Cycler: Perkin Elmer 9700				
Protocol:				
Template: 25 ng				
Primer: each 1 uM				
dNTPs: each 200 uM				
AmpliTaq Gold Polymerase: 0.07 units/ul				
Total Vol: 5 ul				
Buffer:				
MgCl2: 2.5 mM				
KCl: 50 mM				
Tris-HCl: 10 mM				
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REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

Mammalia; Eutheria; Primates; Catarrhini; Hominoidea; Homo.
1 (bases 1 to 177738)
Birren, B., Nussbaum, C. and Lander, E.
Homo sapiens chromosome 17, clone RP11-153A23
Unpublished

2 (bases 1 to 177738)
Birren, B., Linton, L., Nussbaum, C., Lander, E., Abraham, H., Allen, N.,
Anderson, S., Baldwin, J., Barna, N., Beckerly, R., Bida, F.,
Boguslavskiy, L., Boukhalter, B., Brown, A., Burkett, G., Castle, A.,
Chapel, Y., Colangelo, M., Collins, S., Collymore, A., Cooke, P.,
Dearellano, K., Dewar, K., Domino, M., Doyle, M., Fennestor, J.,
Ferrellano, P., FitzHugh, W., Forrest, C., Gage, D., Galagan, J.,
Gardyna, S., Grant, G., Hagos, B., Heatford, A., Horton, L.,
Howland, J., Johnson, R., Jones, C., Kann, L., Karatas, A., Klein, J.,
Lander, T., Lechoczy, J., Levine, R., Liu, C., Liu, G., Locke, K.,
Macdonald, P., Marquis, N., McEwan, P., McGurk, A., McKernan, K.,
McPheters, R., Meldrum, J., Meneus, L., Morrow, J., Naylor, J.,
Norman, C.H., O'Connor, T., O'Donnell, P., Oliver, T.M., Peterson, K.,
Pierre, N., Pisani, C., Pollara, V., Raymond, C., Riley, R., Rothman, D.,
Roy, A., Santos, R., Severy, P., Spencer, B., Stange-Thomann, N.,
Stojanovic, N., Subramanian, A., Talamas, J., Testfaye, S., Theodore, J.,
Tirrell, A., Vassiliev, H., Viel, R., Vo, A., Wu, X., Wyman, D., Ye, W.J.,
Zimmer, A. and Zody, M.
Direct Submission
Submitted (16-JAN-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

3 (bases 1 to 177738)
Birren, B., Nussbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhalter, B.,
Camarata, J., Chang, J., Chararo, B., Choepel, Y., Collymore, A.,
Cooke, A., Cooke, P., Dearellano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,
Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Mihova, T., Mienga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schupback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Testfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (07-JUN-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA

4 (bases 1 to 177738)
Birren, B., Nussbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
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Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gord, S., Graham, L., Grand-Pierre, N., Hagos, B.,
Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C., Kamat, A.,
Karatas, A., Kells, C., Landers, T., Levine, R., Lindblad-Toh, K.,
Liu, G., MacLean, C., Macdonald, P., Major, J., Matthews, C.,
McCarthy, M., Meldrum, J., Meneus, L., Mihova, T., Mienga, V.,
Murphy, T., Naylor, J., Nguyen, C., Nicol, R., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neil, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Raymond, C., Retta, R., Rise, C., Rogov, P.,
Roman, J., Roy, A., Schauer, S., Schupback, R., Seaman, S., Severy, P.,
Smith, C., Spencer, B., Stange-Thomann, N., Stojanovic, N., Talamas, J.,
Testfaye, S., Theodore, J., Topham, K., Travers, M., Vassiliev, H.,
Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.
Direct Submission
Submitted (01-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 1, 2002 this sequence version replaced gi:21702950.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)

http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Web site: http://www-seq.wi.mit.edu
Contact: sequence submissions@genome.wi.mit.edu
----- Project Information
Center project name: L5781
Center clone name: 153_A_23
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RESULT 10
AC087645/c

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DEFINITION AC087645
AC087645.19 GI:27311503
VERSION
KEYWORDS HTG.
SOURCE Homo sapiens (human)

ORGANISM

Homo sapiens
Eukaryota; Euteleostomi;
Mammalia; Eutheria; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

REFERENCE

1 (bases 1 to 209751)

AUTHORS

Birren, B., Nusbaum, C. and Lander, E.

TITLE

Homo sapiens chromosome 17, clone RP11-219G17

JOURNAL

Unpublished

REFERENCE

2 (bases 1 to 209751)

AUTHORS

Birren, B., Linton, L., Nusbaum, C., Lander, E., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Boguslavskiy, L., Boukhgalter, B., Brown, A.,
Camarata, J., Campopiano, A., Choepel, Y., Colangelo, M., Collins, S.,
Collumore, A., Cooke, P., DeArelano, K., Dewar, K., Diaz, J.S.,
Dodge, S., Faro, S., Ferreira, P., FitzHugh, W., Gage, D., Galagan, J.,
Gardyna, S., Ginde, S., Goyette, M., Graham, L., Grand-Pierre, N.,
Hagos, B., Heaford, A., Horton, L., Hulme, W., Iliev, I., Johnson, R.,
Jones, C., Karatas, A., LaRocque, K., Lamazares, R., Landers, T.,
Lehoczky, J., Levine, R., Liu, G., MacLean, C., Macdonald, P.,
Marquis, N., Matthews, C., McCarthy, M., McEwan, P., McKernan, K.,
McPheters, R., Meldrum, J., Meneus, L., Mihova, T., Mienga, V.,
Murphy, T., Naylor, J., Nguyen, C., Norbu, C., Norman, C.H.,
O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J., Peterson, K.,
Phunkhang, P., Pierre, N., Poillara, V., Raymond, C., Retta, R.,
Rieback, M., Riley, R., Rise, C., Rogov, P., Roman, J., Rossetti, M.,
Roy, A., Santos, R., Schauer, S., Schupback, R., Seaman, S., Sever, P.,
Sougniez, C., Spencer, B., Stange-Thomann, N., Stojanovic, N.,
Strauss, N., Subramanian, A., Talamas, J., Testaye, S., Theodore, J.,
Travers, M., Travis, N., Trigilio, J., Vassiliev, H., Viel, R., Vo, A.,
Wilson, B., Wu, X., Wyman, D., Ye, W.J., Young, G., Zainoun, J.,
Zembek, L., Zimmer, A. and Zody, M.

TITLE

Direct Submission

JOURNAL

Submitted (15-JAN-2001) Whitehead Institute/MIT Center for Genome

REFERENCE

Research, 320 Charles Street, Cambridge, MA 02141, USA

AUTHORS

3 (bases 1 to 209751)
Birren, B., Nusbaum, C., Lander, E., Ali, A., Allen, N., Anderson, S.,
Barna, N., Bastien, V., Bloom, T., Boguslavskiy, L., Boukhgalter, B.,
Camarata, J., Chang, J., Chararo, B., Choepel, Y., Collymore, A.,
Cooke, A., Cooke, P., DeArelano, K., Dewar, K., Diaz, J.S., Dodge, S.,
Faro, S., Ferreira, P., FitzGerald, M., Gage, D., Galagan, J.,
Gardyna, S., Gird, S., Graham, L., Grand-Pierre, N., Hafez, N.,
Hagos, B., Horton, L., Hulme, W., Iliev, I., Johnson, R., Jones, C.,
Kamat, A., Karatas, A., Kells, C., Landers, T., Levine, R.,
Lindblad-Toh, K., Liu, G., MacLean, C., Macdonald, P., Major, J.,
Matthews, C., McCarthy, M., Meldrum, J., Meneus, L., Mihova, T.,
Mienga, V., Murphy, T., Naylor, J., Nguyen, C., Nicol, R., North, C.,
Norman, C.H., O'Connor, T., O'Donnell, P., O'Neill, D., Oliver, J.,
Peterson, K., Phunkhang, P., Pierre, N., Raymond, C., Retta, R.,
Rise, C., Rogov, P., Roman, J., Roy, A., Schauer, S., Schupback, R.,
Seaman, S., Sever, P., Smith, C., Spencer, B., Stange-Thomann, N.,
Stojanovic, N., Talamas, J., Testaye, S., Theodore, J., Tophann, K.,
Travers, M., Vassiliev, H., Viel, R., Vo, A., Wilson, B., Wu, X.,
Wyman, D., Young, G., Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.

TITLE

Direct Submission

JOURNAL

Submitted (16-JAN-2003) Whitehead Institute/MIT Center for Genome

REFERENCE

Research, 320 Charles Street, Cambridge, MA 02141, USA

AUTHORS

4 (bases 1 to 209751)
Birren, B., Nusbaum, C., Lander, E., Abouelleil, A., Allen, N.,
Anderson, S., Arachchi, H.M., Barna, N., Bastien, V., Bloom, T.,
Boguslavskiy, L., Boukhgalter, B., Camarata, J., Chang, J., Choepel, Y.,
Collumore, A., Cooke, A., Cooke, P., Corum, B., DeArelano, K.,
Diaz, J.S., Dodge, S., Dooley, K., Dorris, L., Erickson, J., Faro, S.,
Ferreira, P., FitzGerald, M., Gage, D., Galagan, J., Gardyna, S.,
Graham, L., Grand-Pierre, N., Hafez, N., Hagopian, D., Hagos, B.,


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Db 181052 TGGACCAAGACATCAAGTGGCCAAAGAGTGGTGAAGACCCAGCGCGCTCCTT 180993
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Db 180992 CAACCTCAAGCTCACACTCAAGTCCAGTCCAGACCTGCTCCTCACTACTTCGCTGGGCGTC 180933
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RESULT 11
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DEFINITION SAMPLING.
ACCESSION AC010532
VERSION AC010532.2 GI:6758797
KEYWORDS HTG; HTGS PHASE0.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 229426)
AUTHORS DOE Joint Genome Institute.
TITLE Sequencing of Human Chromosome 17
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 229426)
AUTHORS DOE Joint Genome Institute.
TITLE Direct Submission
JOURNAL Submitted (15-SEP-1999) Production Sequencing Facility, DOE Joint
Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
COMMENT On Jan 26, 2000 this sequence version replaced gi:5882406.
-----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
* NOTE: This record contains 99 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
1 563: contig of 563 bp in length
* 1 564 1436: contig of 873 bp in length
* 1437 1649: contig of 213 bp in length
* 1650 2523: contig of 874 bp in length
* 2524 3286: contig of 763 bp in length
* 3287 3613: contig of 327 bp in length
* 3614 5430: contig of 1817 bp in length
* 5431 6681: contig of 1251 bp in length
* 6682 7357: contig of 676 bp in length
* 7358 8117: contig of 760 bp in length
* 8118 8718: contig of 601 bp in length
* 8719 9464: contig of 746 bp in length
* 9465 9964: contig of 500 bp in length
* 9965 10959: contig of 995 bp in length
* 10960 11873: contig of 914 bp in length
* 11874 12656: contig of 783 bp in length
* 12657 13674: contig of 1018 bp in length
* 13675 14935: contig of 1261 bp in length
* 14936 16423: contig of 1488 bp in length
* 16424 17186: contig of 763 bp in length
* 17187 18079: contig of 893 bp in length
* 18080 18813: contig of 734 bp in length
* 18814 19729: contig of 916 bp in length
* 19730 20675: contig of 946 bp in length
* 20676 21721: contig of 1046 bp in length
* 21722 23111: contig of 1390 bp in length
* 23112 25188: contig of 2077 bp in length
* 25189 26588: contig of 1400 bp in length
* 26589 27644: contig of 1056 bp in length
* 27645 29372: contig of 1728 bp in length
* 29373 29495: contig of 123 bp in length
* 29496 30431: contig of 936 bp in length
* 30432 31449: contig of 1018 bp in length
* 31450 32983: contig of 1534 bp in length
* 32984 33162: contig of 179 bp in length
* 33163 33975: contig of 813 bp in length
* 33976 35031: contig of 1056 bp in length
* 35032 35709: contig of 678 bp in length
* 35710 37308: contig of 1599 bp in length
* 37309 38107: contig of 799 bp in length
* 38108 39102: contig of 995 bp in length
* 39103 39591: contig of 489 bp in length
* 39592 41151: contig of 1560 bp in length
* 41152 42842: contig of 1691 bp in length
* 42843 43828: contig of 986 bp in length
* 43829 44009: contig of 181 bp in length
* 44010 44009: contig of 181 bp in length
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Db 135356 CCATCCTTTGCTTCGCGCTCTACAGGAGCACCCCGCTCTCAGTCAAGAGGAGTTTGGGG 135415

QY 808 GGTTCAGGATAGGGAATGGGGAGGTGAGAGACGACGCAAGACGAGCCATGTAGATGAAC 867
Db 135416 GGTTCAGGATAGGGAATGGGGAGGTGAGAGACGACGCAAGACGAGCCATGTAGATGAAC 135475

QY 868 CGTCCAGAGAGCCCAAGCAGGCGAGGACTGCAGGCCATCAGCGTGCACCTGTCGTATTT 927
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QY 928 GGAATTTCATGCAAAATGAGTGTGTTTGTAGTGTCTTTCGCCACAAAAA 987
Db 135536 GGAATTTCATGCAAAATGAGTGTGTTTGTAGTGTCTTTCGCCACAAAAA 135595

QY 988 AAAA 991
Db 135596 AAAA 135599

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ACCESSION  AC032035.3 GI:9994161
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KEYWORDS   Homo sapiens (human)
SOURCE     Homo sapiens
ORGANISM   Homo sapiens
REFERENCE  1 (bases 1 to 154840)
AUTHORS    Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
            Anderson,S., Baldwin,J., Barna,N., Bastien,V., Bada,F.,
            Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G.,
            Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
            Collymore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Gage,D.,
            Dodge,S., Domino,M., Doyle,M., Ferreira,P., Fitzhugh,W., Gage,D.,
            Galagan,J., Gardyna,S., Glnde,S., Goyette,M., Graham,L.,
            Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,
            Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
            Klein,J., LaRoque,K., Lamazares,R., Landers,T., Lehoczy,J.,
            Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
            McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
            Meldrim,J., Meneus,L., Mihova,T., Miranda,C., Mlenga,V., Morrow,J.,
            Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
            O'Neil,D., Olivari,T.M., Oliver,J., Peterson,K., Pierre,N.,
            Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
            Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
            Strange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
            Tesfaye,S., Theodore,J., Tirrell,A., Travers,M., Triggilio,J.,
            Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
            Young,G., Zainoun,J., Zimmer,A. and Zody,M.
            Direct Submission
            Submitted (03-APR-2000) Whitehead Institute/MIT Center for Genome
            Research, 320 Charles Street, Cambridge, MA 02141, USA
            On Sep 8, 2000 this sequence version replaced 91:7705196.
            All repeats were identified using RepeatMasker:
            Smit, A.F.A. & Green, P. (1996-1997)

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http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L9138
 Center clone name: 141_D_15

* NOTE: This is a 'working draft' sequence. It currently
 * consists of 18 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

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 * 7316 9541: contig of 2226 bp in length
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 * 9642 12211: contig of 2570 bp in length
 * 12212 12311: gap of 100 bp
 * 12312 14923: contig of 2612 bp in length
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 * 19095 19194: gap of 100 bp
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 * 22387 26147: contig of 3761 bp in length
 * 26148 26247: gap of 100 bp
 * 26248 31933: contig of 5686 bp in length
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 * 37340 37439: gap of 100 bp
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 * 42959 48810: contig of 5852 bp in length
 * 48811 48910: gap of 100 bp
 * 48911 56029: contig of 7119 bp in length
 * 56030 56129: gap of 100 bp
 * 56130 62189: contig of 6060 bp in length
 * 62190 62289: gap of 100 bp
 * 62290 82959: contig of 20670 bp in length
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 * 83060 107059: contig of 24000 bp in length
 * 107060 107159: gap of 100 bp
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FEATURES

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 /chromosome="17"
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 Best Local Similarity 97.4%; Pred. No. 1.7e-59;
 Matches 296; Conservative 0; Mismatches 8; Indels 0; Gaps 0;
 QY 688 CAGTGTGCCCCCAGCTGTGTACCAAGATGGAGACTGGAGGTCCTCCCTGGAGAGCC 747
 |||||
 Db 86135 CACTGCTTTGTCAAGGTGTGTACCAAGATGGAGACTGGAGGTCCTCCCTGGAGAGCC 86194

QY 748 CCATCCTTCCTTGGCGCTCTACAGAGCACCCGCGCTCTGAGTGAAGAGGATTTGGGG 807
 Db 86195 CCATCCTTTCCTTGGCGCTCTACAGAGCACCCGCGCTTGTAGTGAAGAGGATTTGGGG 86254
 QY 808 GCTTCAGATAGGGAATGGGAGGTTCAGAGGACGCAAAAGCAGCAGCATGTAGATGAAC 867
 Db 86255 GTTTCAGATAGGGAATGGGAGGTTCAGAGGACGCAAAAGCAGCAGCATGTAGATGAAC 86314
 QY 868 COTTCAGAGAGCCAAAGCAGCGGAGGACTGAGGCGCATCAGGCGCATGTTCTGATTT 927
 Db 86315 COTTCAGAGAGCCAAAGCAGCGGAGGACTGAGGCGCATCAGGCGCATGTTCTGATTT 86374
 QY 928 GGAGTTTCATGCAAAATGAGTGTGTTTACCTCTTGCACAAAAAATAAAAAA 987
 Db 86375 GGAGTTTCATGCAAAATGAGTGTGTTTACCTCTTGCACAAAAAATAAAAAA 86434
 QY 988 AAAA 991
 Db 86435 AAAA 86438
 RESULT 13
 AC010532/c
 LOCUS
 DEFINITION Homo sapiens chromosome 17 clone RP11-219G17, LOW-PASS SEQUENCE
 SAMPLING.
 AC010532
 VERSION AC010532.2 GI:6758797
 KEYWORDS HTG; HTGS_PHASE0.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE
 1 (bases 1 to 229426)
 AUTHORS DOE Joint Genome Institute.
 TITLE Sequencing of Human Chromosome 17
 JOURNAL Unpublished
 REFERENCE 2 (bases 1 to 229426)
 AUTHORS DOE Joint Genome Institute.
 TITLE Direct SubtilBion
 JOURNAL Submitted (15-SEP-1999) Production Sequencing Facility, DOE Joint
 Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
 COMMENT On Jan 26, 2000 this sequence version replaced gi:5882406.
 -----Genome Center
 Center: Joint Genome Institute
 Center Code: JGI
 Web site: http://www.jgi.doe.gov

 * NOTE: This record contains 99 individual
 * sequencing reads that have not been assembled into
 * contigs. Runs of N are used to separate the reads
 * and the order in which they appear is completely
 * arbitrary. Low-pass sequence sampling is useful for
 * identifying clones that may be gene-rich and allows
 * overlap relationships among clones to be deduced.
 * However, it should not be assumed that this clone
 * will be sequenced to completion. In the event that
 * the record is updated, the accession number will
 * be preserved.
 * 1 563: contig of 563 bp in length
 * gap of unknown length
 * 564 1436: contig of 873 bp in length
 * gap of unknown length
 * 1437 1649: contig of 213 bp in length
 * gap of unknown length
 * 1650 2523: contig of 874 bp in length
 * gap of unknown length
 * 2524 3286: contig of 763 bp in length
 * gap of unknown length
 * 3287 3613: contig of 327 bp in length
 * gap of unknown length
 * 3614 5430: contig of 1817 bp in length
 * gap of unknown length

* 5431	6681:	contig of 1251 bp in length	* 42843	43828:	gap of unknown length
* 6682	7357:	gap of unknown length	* 43829	44009:	contig of 986 bp in length
* 7358	8117:	gap of unknown length	* 44010	44874:	gap of unknown length
* 8118	8718:	contig of 601 bp in length	* 44875	45191:	contig of 865 bp in length
* 8719	9464:	gap of unknown length	* 45192	46147:	gap of unknown length
* 9465	9964:	contig of 746 bp in length	* 46148	47504:	gap of unknown length
* 9965	10959:	gap of unknown length	* 47505	49091:	contig of 956 bp in length
* 10960	11873:	contig of 995 bp in length	* 49092	50259:	gap of unknown length
* 11874	12656:	gap of unknown length	* 50260	51361:	contig of 1168 bp in length
* 12657	13674:	gap of unknown length	* 51362	52631:	gap of unknown length
* 13675	14935:	contig of 1018 bp in length	* 52632	53607:	contig of 1102 bp in length
* 14936	16423:	gap of unknown length	* 53608	54820:	contig of 1270 bp in length
* 16424	17186:	contig of 1488 bp in length	* 54821	56600:	gap of unknown length
* 17187	18079:	gap of unknown length	* 56601	58236:	contig of 1780 bp in length
* 18080	18813:	gap of unknown length	* 58237	59716:	contig of 1636 bp in length
* 18814	19729:	contig of 734 bp in length	* 59717	62221:	gap of unknown length
* 19730	20675:	gap of unknown length	* 62222	65085:	contig of 1480 bp in length
* 20676	21721:	contig of 916 bp in length	* 65086	66739:	gap of unknown length
* 21722	23111:	gap of unknown length	* 66740	68359:	contig of 1654 bp in length
* 23112	25188:	contig of 1390 bp in length	* 68360	70014:	gap of unknown length
* 25189	26588:	gap of unknown length	* 70015	71808:	contig of 1620 bp in length
* 26589	27644:	gap of unknown length	* 71809	73581:	gap of unknown length
* 27645	29372:	contig of 1056 bp in length	* 73582	75574:	contig of 1794 bp in length
* 29373	29495:	gap of unknown length	* 75575	78468:	gap of unknown length
* 29496	30431:	contig of 123 bp in length	* 78469	79231:	gap of unknown length
* 30432	31449:	gap of unknown length	* 79232	83240:	contig of 763 bp in length
* 31450	32983:	contig of 1018 bp in length	* 83241	86157:	gap of unknown length
* 32984	33162:	gap of unknown length	* 86158	89150:	contig of 4009 bp in length
* 33163	33975:	contig of 1534 bp in length	* 89151	92671:	gap of unknown length
* 33976	35031:	gap of unknown length	* 92672	95442:	contig of 3521 bp in length
* 35032	35709:	gap of unknown length	* 95443	98476:	gap of unknown length
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* 37309	38107:	gap of unknown length	* 100816	103005:	gap of unknown length
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* 39592	41151:	contig of 489 bp in length	* 108705	111097:	contig of 3190 bp in length
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*      gap of unknown length
* 114889 117423: contig of 2535 bp in length
*      gap of unknown length

Query Match
Best Local Similarity 18.1%; Score 179.4; DB 2; Length 229426;
Matches 183; Conservative 0; Mismatches 6; Indels 0; Gaps 0;

QY 244 CCTCTGTGGAACCAAGATCAATCAAGTGGCCAAAGAGGTGGTGAAGACCCACGAGCGG 303
Db 1331 CGTTCGGGGAACCCCAAGATCAATCAAGTGGCCAAAGAGGTGGTGAAGACCCACGAGCGG 1272
QY 304 CTTCTTCAACCTCAAGTCACTCACTCAAGTCCAGTCCAGACCTGCTCACTACTTTGCGC 363
Db 1271 CTTCTTCAACCTCAAGTCACTCACTCAAGTCCAGTCCAGACCTGCTCACTACTTTGCT 1212
QY 364 GGGGCTCTCCACCTCAAGTGGCCATGTGGACAGTGCAGGCTACAGATGCATGCGGAGC 423
Db 1211 GGGGCTCTCCACCTCAAGTGGCCATGTGGACAGTGCAGGCTACAGATGCATGCGGAGC 423
QY 424 TGTGTGTCCA 432
Db 1151 TGTGTGTCCA 1143

RESULT 14
AL645856
LOCUS
DEFINITION
  Mouse DNA sequence from clone RP23-151N19 on chromosome 11,
  complete sequence.
ACCESSION
  AL645856
VERSION
  AL645856.5 GI:22531400
KEYWORDS
  HTG.
SOURCE
  Mus musculus (house mouse)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
  1 (bases 1 to 116300)

REFERENCE
  1
  Submitted (24-AUG-2002) Wellcome Trust Sanger Institute, Hinxton,
  Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
  humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
  On Aug 27, 2002 this sequence version replaced gi:21739435.
  ----- Genome Center
  Center: Wellcome Trust Sanger Institute
  Center code: SC
  Web site: http://www.sanger.ac.uk
  Contact: humquery@sanger.ac.uk

```

During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TREMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep RP23-151N19 is from the RP23-151 Mouse PAC Library

constructed by the group of Pieter de Jong.

For further details see http://www.chori.org/bacpac/home.htm

VECTOR: pBac3.6.

```

FEATURES
  source
  Location/Qualifiers
    1..116300
      /organism="Mus musculus"
      /mol_type="genomic DNA"
      /db_xref="taxon:10090"
      /chromosome="11"
      /clone="RP23-151N19"
      /clone_lib="RPCI-23"
BASE COUNT 29288 28679 C 29788 G 28545 T
ORIGIN

Query Match
Best Local Similarity 16.5%; Score 163.4; DB 10; Length 116300;
Matches 209; Conservative 0; Mismatches 76; Indels 0; Gaps 0;

QY 148 TCTCCATTGCTCAACAAGTCTCTGGAAGTTTCCCAAGAGCGCGTGGTGTCTATAACCT 207
Db 93425 TCACCATGCTCAACAAGTACTGGAAGTTTATCCCAAGCGGAGGTGCTTTATAACCT 93484
QY 208 GCTGTGCACCCCAAGCCACCCATCATCTATTTCCCTCTGTGGAAACCAAGAACATCA 267
Db 93485 GCGATGCCCTGAGCGGTCCAGCCCATCACATCTCTCTGGCTAGCGGAGTATCC 93544
QY 268 AGGTGCCAAGAAGGTGGTGAAGACCCAGAGCGCGCTCTTCAACCTCAACGTCAAC 327
Db 93545 TGGTGGCAAAAAGGTTGTGATGACTCGGTGCGCGCTCTTCAACATCAATATCA 93604
QY 328 TCAAGTCCAGTCCAGACCTGCTCACTACTTCTGCGGGGCTCTCCACCTCAAGTGC 387
Db 93605 TCAAGTCCAGCCAGACCTGCTCACTACTTCTGCGGAGCACTCGAATCTGGCACCT 93664
QY 388 ATGTGGACAGTGCAGGCTACAGATGACATGCGGAGCTGTGGTCCA 432
Db 93665 ATGGACCCAGCAGAGCTCCAGATGTACAGAACTGTGGGCTA 93709

RESULT 15
AL954690
LOCUS
DEFINITION
  Mus musculus chromosome 11 clone RP23-386A4, linear
  HTG 15-NOV-2002
  PROGRESS ***
ACCESSION
  AL954690
VERSION
  AL954690.2 GI:25136861
KEYWORDS
  HTG; HTGS PHASE2; HTGS CANCELLED.
SOURCE
  Mus musculus (house mouse)
ORGANISM
  Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
  Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
  1 (bases 1 to 197194)

REFERENCE
  1
  Submitted (13-NOV-2002) Wellcome Trust Sanger Institute, Hinxton,
  Cambridgeshire, CB10 1SA, UK. E-mail enquiries:
  humquery@sanger.ac.uk Clone requests: clonerequest@sanger.ac.uk
  On Nov 19, 2002 this sequence version replaced gi:25045706.
  Sequence from the Mouse Genome Sequencing Consortium whole genome
  shotgun may have been used to confirm this sequence. Sequence data
  from the whole genome shotgun alone has only been used where it has
  a phred quality of at least 30.
  ----- Genome Center
  Center: Wellcome Trust Sanger Institute
  Center code: SC
  Web site: http://www.sanger.ac.uk
  Contact: humquery@sanger.ac.uk
  ----- Project Information
  Center project name: BM386A4
  ----- Summary Statistics
  Assembly program: XGAP4; version 4.5
  Chemistry: Dye-terminator; 100% of reads
  Consensus quality: 195990 bases at least Q40
  Consensus quality: 196234 bases at least Q30
  Consensus quality: 196338 bases at least Q20
  Insert size: 197194; sum-of-contigs

```

```

Insert size: 189681; 3.6% error; agarose-fp
Quality coverage: 7.24x in Q20 bases; sum-of-contigs Quality
coverage: 7.53x in Q20 bases; agarose-fp
-----
* NOTE: This is a 'working draft' sequence.
* This sequence will be replaced
* by the finished sequence as soon as it is available and
* the accession number will be preserved.
FEATURES
    source
        1..197194
            /organism="Mus musculus"
            /mol_type="genomic DNA"
            /db_xref="taxon:10090"
            /chromosome="11"
            /clone="RP23-386A4"
            /clone_lib="RPCI-23"
            /note="assembly_fragment:02398"
    misc_feature
        1..197194
            clone_end:SP6
            vector_side:left"
BASE COUNT  46441 a 51277 c 51850 g 47922 t      4 others
ORIGIN
Query Match      16.5%; Score 163.4; DB 2; Length 197194;
Best Local Similarity 73.3%; Pred. No. 1.1e-28;
Matches 209; Conservative 0; Mismatches 76; Indels 0; Gaps 0;
QY 148 TCTCATTCCTTACAAAGTCCTGGAGTTTCCCAAGGCGGTGGTGTCTCAACCT 207
Db 20120 TCACCATTCCTTACAAAGTCTGGAAGTTTCCCAAGGCGGAGGTGCTTAACT 20179
QY 208 GCTGTGACCCAGCACCACCGCCATCCTATTCCTCTGTGGACCAAGACATCA 267
Db 20180 GCGATGCCCCCTGAGCGCTCCAGCCCATCACATCTCTCTCTGCTAGCGAGTATCC 20239
QY 268 AGTTGGCCCAAGAGTGTGGAAGACCCAGCGCGGCTCTTCAACTCAACGTCACAC 327
Db 20240 TGTGGCAAAAAGTGTGATGACTCGTGCGCGGCTCTTCAACATCATATCACA 20299
QY 328 TCAAGTCAGTCCAGACTGTCTACCTACTTGTCCGGCGGCTCTCCACCTCAGTGCCC 387
Db 20300 TCAAGTCAGCCAGACCTGTCTACCTACTCTGCGAGGCAACCTCGAACCTTGACACCT 20359
QY 388 ATGTGGACAGTCCAGGCTACAGATGACACTGGAGCTGTGTCCA 432
Db 20360 ATGGACCCAGCAGCAGGCTCCAGATGTACCGAAGTGTGGCTA 20404

RESULT 16
AC100565
LOCUS      AC100565      49532 bp      DNA      linear      HTG 22-NOV-2001
DEFINITION Mus musculus clone RP23-154K21, LOW-PASS SEQUENCE SAMPLING.
ACCESSION  AC100565
VERSION    AC100565.1 GI:17047931
KEYWORDS  HTG; HTGS PHASE0.
SOURCE    Mus musculus (house mouse)
ORGANISM  Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 49532)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
TITLE      Mus musculus, clone RP23-154K21
JOURNAL    Unpublished
REFERENCE  2 (bases 1 to 49532)
AUTHORS    Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguelavkiy,L., Boukhalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepei,Y., Colangelo,M., Collins,S., Collymore,A., Cook,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
Gardos,B., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,

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Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrim,J.,
Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,S., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Traversan,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (22-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L15700
Center clone name: 154_K_21
-----
* NOTE: This record contains 48 individual
* sequencing reads that have not been assembled into
* contigs. Runs of N are used to separate the reads
* and the order in which they appear is completely
* arbitrary. Low-pass sequence sampling is useful for
* identifying clones that may be gene-rich and allows
* overlap relationships among clones to be deduced.
* However, it should not be assumed that this clone
* will be sequenced to completion. In the event that
* the record is updated, the accession number will
* be preserved.
1 986: contig of 986 bp in length
* 987 1086: gap of 100 bp
* 1087 2009: contig of 923 bp in length
* 2010 2109: gap of 100 bp
* 2110 3007: contig of 898 bp in length
* 3008 3107: gap of 100 bp
* 3108 3866: contig of 759 bp in length
* 3867 3966: gap of 100 bp
* 3967 4874: contig of 908 bp in length
* 4875 4974: gap of 100 bp
* 4975 5839: contig of 865 bp in length
* 5840 5939: gap of 100 bp
* 5940 7017: contig of 1078 bp in length
* 7018 7117: gap of 100 bp
* 7118 8079: contig of 962 bp in length
* 8080 8179: gap of 100 bp
* 8180 9139: contig of 960 bp in length
* 9140 9239: gap of 100 bp
* 9240 10254: contig of 1015 bp in length
* 10255 10354: gap of 100 bp
* 10355 11289: contig of 935 bp in length
* 11290 11389: gap of 100 bp
* 11390 12334: contig of 945 bp in length
* 12335 12435: gap of 100 bp
* 12435 13410: contig of 976 bp in length
* 13411 13510: gap of 100 bp
* 13511 14445: contig of 935 bp in length
* 14446 14545: gap of 100 bp
* 14546 15272: contig of 727 bp in length
* 15273 15373: gap of 100 bp
* 15373 16394: contig of 1022 bp in length
* 16395 16494: gap of 100 bp
* 16495 17461: contig of 967 bp in length
* 17462 17561: gap of 100 bp

```

TITLE
JOURNAL

COMMENT


```

Mg2+: 1.5 mM
KCl: 50 mM
Tris-HCl: 10 mM
Gelatin: .001 %
Location/Qualifiers
1. .150
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="Human THudson EST"
/note="STS derived from sequences in dbEST and the
Unigene collection."
1. .150
1. .20
primer_bind complement(132..150)
primer_bind 30 a 46 c 33 g 40 t 1 others
BASE COUNT
ORIGIN
Query Match 15.0%; Score 149; DB 11; Length 150;
Best Local Similarity 99.3%; Pred. No. 3.8e-25;
Matches 149; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 820 GGAATGGGAGGTGACAGGAGCGCAAGCAGGAGCATGTAAGTAAACCCGCCAGAGGC 879
Db 150 GGAATGGGAGGTGACAGGAGCGCAAGCAGGAGCATGTAAGTAAACCCGCCAGAGGC 91
Qy 880 CAAGCAGCGCAGAGACTGCAGGCGATCAGCGTGCACCTGTCGATTTGGAGTTCATGCA 939
Db 90 CAAGCAGCGCAGAGACTGCAGGCGATCAGCGTGCACCTGTCGATTTGGAGTTCATGCA 31
Qy 940 AAATGAGTGTGTTTAGTCTCTTGGCCAC 969
Db 30 AAATGAGTGTGTTTAGTCTCTTGGCCAC 1

RESULT 18
G43546/c
LOCUS
DEFINITION
WIAF-2407-STS Human THudson EST Homo sapiens STS cDNA, sequence
tagged site.
G43546
ACCESSION
VERSION G43546.1 GI:14192463
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominiidae; Homo.
1 (bases 1 to 150)
Wang, D.G., Fan, J.B., Siao, C.J., Berno, A., Young, P., Sapolsky, R.,
Ghandour, G., Perkins, N., Winchester, E., Spencer, J., Kruglyak, L.,
Stein, L., Hsie, L., Topaloglou, T., Hubbell, E., Robinson, E.,
Mittmann, M., Morris, M.S., Shen, N., Kilburn, D., Rioux, J.,
Nusbaum, C., Rozen, S., Hudson, T.J., Lipschutz, R., Chee, M. and
Lander, E.S.
Large-scale identification, mapping, and genotyping of
single-nucleotide polymorphisms in the human genome
Science 280 (5366), 1077-1082 (1998)
98248615
9582121
Synonyms: EST376246b, EST376246
Contact: Thomas Hudson
Whitehead Institute/MIT Center for Genome Research
Whitehead Institute for Biomedical Research
9 Cambridge Center, Cambridge MA 02142 USA
Tel: 617 252 1900
Fax: 617 252 1902
Email: thudson@genome.wi.mit.edu
Primer A: GTGGCAAGCAGCTAAAC
Primer B: GGAATGGGAGGTGACAGG
STS size: 150
PCR Profile:
Presoak: 94 degrees C for 4.00 minutes
Denaturation: 94 degrees C for 50.0 seconds

```

```

Annealing: 58 degrees C for 1.50 minutes
Polymerization: 72 degrees C for 1.00 minutes
PCR Cycles: 30
Thermal Cycler: custom built by IAS, Costar, Cambridge MA

Protocol:
Template: 10 ng
Primer: each 5 pM
dNTPs: 4 mM
Taq Polymerase: 0.5 U
Total Vol: 20 uL

Buffer:
Mg2+: 1.5 mM
KCl: 50 mM
Tris-HCl: 10 mM
Gelatin: .001 %
Location/Qualifiers
1. .150
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone_lib="Human THudson EST"
/note="STS derived from sequences in dbEST and the
Unigene collection."
1. .150
1. .20
primer_bind complement(132..150)
primer_bind 30 a 47 c 32 g 40 t 1 others
BASE COUNT
ORIGIN
Query Match 15.0%; Score 149; DB 11; Length 150;
Best Local Similarity 99.3%; Pred. No. 3.8e-25;
Matches 149; Conservative 0; Mismatches 1; Indels 0; Gaps 0;

Qy 820 GGAATGGGAGGTGACAGGAGCGCAAGCAGGAGCATGTAAGTAAACCCGCCAGAGGC 879
Db 150 GGAATGGGAGGTGACAGGAGCGCAAGCAGGAGCATGTAAGTAAACCCGCCAGAGGC 91
Qy 880 CAAGCAGCGCAGAGACTGCAGGCGATCAGCGTGCACCTGTCGATTTGGAGTTCATGCA 939
Db 90 CAAGCAGCGCAGAGACTGCAGGCGATCAGCGTGCACCTGTCGATTTGGAGTTCATGCA 31
Qy 940 AAATGAGTGTGTTTAGTCTCTTGGCCAC 969
Db 30 AAATGAGTGTGTTTAGTCTCTTGGCCAC 1

RESULT 19
AC143361
LOCUS
DEFINITION
Macaca mulatta clone CH250-271N12, *** SEQUENCING IN PROGRESS ***.
ACCESSION
VERSION AC143361.1 GI:29648548
KEYWORDS
HTG; HTGS PHASE2; HTGS PGI.
SOURCE
Macaca mulatta
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Cercopitheidae;
Cercopitheciinae; Macaca.
1 (bases 1 to 161043)
Csuros, M. and Milosavljevic, A.
Pooled genomic indexing (PGI): mathematical analysis and experiment
design
(in) Guigo, R. and Gusfield, D. (Eds.);
ALGORITHMS IN BIOINFORMATICS, SECOND INTERNATIONAL WORKSHOP, WABI
2002, ROME, ITALY, SEPTEMBER 17-21, 2002, PROCEEDINGS: 10-28;
Springer (2002)
2 (bases 1 to 161043)
Milosavljevic, A., Sodergren, E., Csuros, M., Li, B., Jackson, A.R.,
Adams, C., Adio-Oduola, B., Ali-ouman, F.R., Allen, C., Alsbrooks, S.L.,
Anaratunge, H.C., Are, J.R., Ayele, M., Banks, T., Barbieri, J.,
Benton, J., Bimaga, K., Blankenburg, K., Bonnin, D., Bouck, J.,
Bowie, S., Brieva, M., Brown, E., Brown, M., Bryant, N.P., Buhay, C.,

```


* the accession number will be preserved.	
FEATURES	* 1 161043: contig of 161043 bp in length.
source	Location/Qualifiers
	1..161043
	/organism="Macaca mulatta"
	/mol_type="genomic DNA"
	/db_xref="taxon:9544"
	/clone="CH250-271N12"
misc_feature	1..161043
	/note="assembly_name:CH250-271N12.1B
	CONFIDENCE: 0.83
BASE COUNT	2845 a 3627 c 3686 g 2810 t 148075 others
ORIGIN	
	Query Match 12.9%; Score 127.8; DB 2; Length 161043;
	Best Local Similarity 76.3%; Pred. No. 4.2e-20;
	Matches 135; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
QY	529 CTATCACCAACAGCCTGATCTGGGAAGGATGGGCAGGTCCACCTGCAGCAGACACCATGCC 588
Db	1 CGANNAGTTCATCTCCAGCAGCGACACGCC 60
QY	589 ACAGCGAGCGCTGCCAATCTCTCTCTCTCGCAGCAGCAGACATCGGACTGGTTCGTGGTCC 648
Db	61 ACGGGCAGCGCTGCCAATCTCTCTCTCTCGCGGGCAGACATCGGACTGGTTCAGTGGC 120
QY	649 AGGCTGCAAAACAGCGCATGTCAGCAGCAGCGCCTCACAGTGGTGCCGCCAGGTG 705
Db	121 AGGCTGCAAAACAGCGTCAATGTCAGCAGCGCCTCACATGGTGCCGCCAGGTG 177
RESULT 20	
AC003666	
LOCUS	151750 bp DNA linear PRI 07-AUG-1999
DEFINITION	Homo sapiens Xp22 BAC GS-551019 (Genome Systems Human BAC library) and cosmids U199A7 and U209F2 (Lawrence Livermore X chromosome cosmid library) containing part of human chloride channel 4 gene, complete sequence.
ACCESSION	AC003666
VERSION	AC002358
KEYWORDS	AC002360 AC003017
SOURCE	GI:2992476
ORGANISM	Homo sapiens (human)
REFERENCE	
AUTHORS	Muzny, D., Aronson, A.D., Brundage, E., Carvelli, K., Chen, E., Chen, J., Di, W., Ding, Y., Dugan, S., Durbin, J., Forcum, J., Ganesh, R., Garcia, C., Goodman, M., Gorrell, J.H., Haywood, M., Jackson, L., Jin, S., Kampa, R., Karpathy, S., Leal, B., Li, Y., Liu, W., Logan, O., Lu, J., Ly, T., Martinez, C., Oswal, G., Perez, L., Rashid, N.D., Rowland, K., Savage, L., Scherer, S.S., Shen, H., Timms, K.M., Todd, J., Vo, O., Worley, K.C., Yu, W., Chinault, C., Nelson, D. and Gibbs, R.A.
TITLE	Direct Submission
JOURNAL	Unpublished
REFERENCE	2 (bases 1 to 151750)
AUTHORS	Worley, K.C.
TITLE	Direct Submission
JOURNAL	Submitted (09-DEC-1997) Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	3 (bases 1 to 151750)
AUTHORS	Worley, K.C.
TITLE	Direct Submission
JOURNAL	Submitted (27-MAR-1998) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	4 (bases 1 to 151750)
AUTHORS	Worley, K.C.
TITLE	Direct Submission
JOURNAL	Submitted (05-MAY-1998) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA
REFERENCE	5 (bases 1 to 151750)

AUTHORS
TITLE
JOURNAL

Worley, K.C.
Direct Submission
Submitted (30-JUL-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
6 (bases 1 to 151750)

REFERENCE
AUTHORS
TITLE
JOURNAL

Worley, K.C.
Direct Submission
Submitted (07-AUG-1999) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
Sequencing is completed to a minimum standard of double strand
coverage with a minimum of 2 clones and 2 reads with no ambiguities
or 2 chemistries with a minimum of 2 clones and 3 reads with no
ambiguities. If the sequence quality does not meet this standard,
it will be indicated in the annotation.

COMMENT

The repeat regions shown were identified using RepeatMasker by
Adrian Smit.

Sequence similarities were identified using PowerBlast by Jinghui
Zhang.

Exon/Intron boundaries of identified genes were chosen if there
were canonical splice junctions that maintained sequence continuity
across the splice junctions.

FEATURES
source

1..151750
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="X"
/map="Xp22"
/clone="GS-551019, U199A7, U209F2"
/clone_lib="Genome Systems Human BAC library, Lawrence
Livermore X chromosome library"

misc_feature

1..374
/note="Overlap bases 16683 to 167256 in AC008008"
/function="Overlap with clone AC008008"

repeat_region

complement(2657..2689)

repeat_region

/rpt_family="AT rich"

repeat_region

complement(2773..2863)

repeat_region

/rpt_family="MERSE"

repeat_region

3472..4871

repeat_region

/rpt_family="L1PA2"

repeat_region

5843..6143

repeat_region

/rpt_family="L1MA8"

repeat_region

complement(6989..7493)

repeat_region

/rpt_family="L1MA3"

repeat_region

complement(7513..7811)

repeat_region

/rpt_family="AluSx"

STS

11128..11201

repeat_region

/rpt_family="L1PA2"

repeat_region

11487..11618

repeat_region

/rpt_family="L1PA2"

repeat_region

/standard_name="hz-1"

repeat_region

11757..11981

repeat_region

/rpt_family="MER20"

repeat_region

12660..12782

repeat_region

/rpt_family="L2"

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

repeat_region

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repeat_region

repeat_region

STS

36178..38428

STS

36898..37069

STS

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29209..29310

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30127..30683

/rpt_family="MLTID"

31072..31160

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complement(33455..33479)

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/rpt_family="MIR"
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Best Local Similarity 7.0%; Score 69.2; DB 9; Length 151750;
Matches 131; Conservative 0; Mismatches 58; Indels 7; Gaps 2;

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Db |||||
QY 5846 AGGAGTTAACAGAGGGTGGATAGGAGTGGAGTGGAGTACAGAGCAGCGTA 5905
Db |||||
QY 856 TGTAGATGAACCGTCCAGAGAGCAAGCAGCGCA---GAGSACTGCAGGCCATCAGCGT 912
Db |||||
QY 5906 TGTAGATGAACAAATCTGGAGATCTAACACACACACCTGAGGACTACATGGCATATAAATT 5965
Db |||||
QY 913 GCACGTGTCGTATTTGGAGTTCATGCAAAAGCAGTGTCTTTAGTGTCTCTGCACAAA 972
Db |||||
QY 5966 ATACTGA---ATTGGGATTCATAAGAAATGAGTAGATTTTAGTGTCTCTGCACAAA 6021
Db |||||
QY 973 AAAAAAAAAAAAAAAAAA 988
Db |||||
QY 6022 AATAAAGCAAAAAA 6037
Db |||||

RESULT 21
AC103588 208670 bp DNA linear PRI 21-MAR-2002
LOCUS Homo sapiens chromosome 3 clone RP11-640H5, complete sequence.
DEFINITION AC103588
ACCESSION AC103588
VERSION AC103588.2 GI:19570161
KEYWORDS HTG.
SOURCE Homo sapiens
ORGANISM Homo sapiens (human)
REFERENCE 1 (bases 1 to 208670)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 208670)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimmachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
Direct Submission
Unpublished
2 (bases 1 to 208670)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Direct Submission
Submitted (29-NOV-2001) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
3 (bases 1 to 208670)
AUTHORS Kaul,R.K., Olson,M.V., Zhou,Y., James,R.A., Rouse,G., Wu,Z.,
Saenphimmachak,C., Phelps,K.A., Raymond,C. and Haugen,E.D.
Direct Submission
Submitted (21-MAR-2002) Genome Center, University of Washington,
Box 352145, Seattle, WA 98195, USA
On Mar 21, 2002 this sequence version replaced gi:17149452.
COMMENT ----- Genome Center
Center: University of Washington Genome Center
Center Code: UWGC
Web site: http://www.genome.washington.edu
Contact: uwgchgs@u.washington.edu
----- Project Information
Center project name: chr-3
Center clone name: RP11-640H5 (bc0507)
----- Summary Statistics

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Sequencing vector: plasmid; 108752; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.990319
 Consensus quality: 208594 bases at least Q40
 Consensus quality: 208666 bases at least Q30
 Consensus quality: 208670 bases at least Q20
 Insert size: 207271; sum-of-contigs
 Quality coverage: 7.7x in Q20 bases; sum-of-contigs

Overlapping Sequences:

5': Mapping in progress
 3': RP11-1C5 (UWGC:bc0101) AC093411, 107464-bp overlap

Sequence Quality Assessment:

This entry has been annotated with sequence quality estimates computed by the Phrap assembly program. All manually edited bases have been reduced to quality zero. Quality levels above 40 are expected to have less than 1 error in 10,000 bp.
 Base-by-base quality values are not generally visible from the GenBank flat file format but are available as part of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., Phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest fingerprinting. Comparison of the experimentally derived digest fragments with sequence-predicted fragments is given below. The electronically-digested sequence consists of both insert and vector, in order to accurately represent the entire circular BAC. Small fragments below a variable cutoff (approximately 400-800 bp) are not resolved in the fingerprint and hence do not appear in the table. There are no significant remaining discrepancies between the experimental and predicted values. Uniquely ordered fragments are separated by dashed lines.

EcoRI				BglII				HindIII			
SeqDerMap	EngPrnt	SeqDerMap	EngPrnt	SeqDerMap	EngPrnt	SeqDerMap	EngPrnt	SeqDerMap	EngPrnt	SeqDerMap	EngPrnt
8696	8812	2524	2527	561	<800	2527	561	<800			
6	<800	2067	2075	6382	6445						
1516	1544	9401	9533	512	<800						
5561	5606	17986	18248	449	<800						
10971	10860	4692	4675	2484	2483						
2516	2558	2347	2350	11918	12069						
1834	1834	7017	7123	511	<800						
2909	2978	3961	3953	1038	1087						
1008	1008	1756	1735	4818	4777						
2704	2761	6058	5990	1677	1673						
13906	13758	5740	5669	2657	2610						
1875	1834	4526	4503	1348	1312						
3723	3725	3178	3199	462	<800						

abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Wp:, WORMPEP; Information on the WORMPEP database can be found at http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping Group. Further information can be found at <http://www.sanger.ac.uk/HGP/Chr9> RP11-497K15 is from the library RP11-11.2 constructed by the group of Pieter de Jong. For further details see <http://www.chori.org/bacpac/home.htm>

VFC10R: pBAC3.6
This sequence is the entire insert of clone RP11-497K15 The true left end of clone RP11-40C6 is at 128837 in this sequence. The true right end of clone RP11-5802 is at 80971 in this sequence.

FEATURES

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	7. .187	/organism="Homo sapiens"	16114. .16288
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	507. .633	/db_xref="taxon:9606"	16646. .16717
	649. .751	/chromosome="9"	/note="MLT1J repeat: matches 104. .187 of consensus"
	133. .1428	/clone="RP11-497K15"	16802. .17084
	3216. .3644	/clone_lib="RP11-11.2"	/note="L1MB8 repeat: matches 5878. .6173 of consensus"
	6009. .6477		17504. .17660
	6702. .6982		/note="L1MA6 repeat: matches 5296. .5456 of consensus"
	7119. .7203		17657. .18355
	7337. .8076		/note="L1MB3 repeat: matches 5349. .6046 of consensus"
	8330. .8613		18357. .18656
	8633. .8927		/note="AluSp repeat: matches 1. .302 of consensus"
	9107. .9266		19964. .20259
	9700. .10001		/note="AluSq repeat: matches 5. .299 of consensus"
	10683. .10783		20604. .20912
	10890. .10969		/note="L2 repeat: matches 302. .628 of consensus"
	11931. .11992		20983. .21292
	12476. .12860		/note="AluY repeat: matches 1. .309 of consensus"
	13210. .13318		21321. .21428
	13898. .14064		/note="FLAM A repeat: matches 21. .128 of consensus"
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	16037. .16512		/note="MIR repeat: matches 30. .125 of consensus"
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			33717. .34022
			/note="AluSc repeat: matches 1. .306 of consensus"
			34170. .34419
			/note="MLT1G repeat: matches 31. .260 of consensus"
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			35085. .35128
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			35525. .35836
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			35844. .35911
			/note="MLT1I repeat: matches 349. .409 of consensus"
			36507. .36760
			/note="L1PB2 repeat: matches 4128. .4385 of consensus"
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			38795. .39098
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Matches 117; Conservative 0; Mismatches 72; Indels 1; Gaps 1;

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QY 862 ATGACCGTCCAGAGCCAGCAGCGGAGGACTGCGGCCATCAGCGTGCATCTTC 921
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Db 76629 ATGAACAAAACTAGAAGACTAATGTACACATGATTTCTAGAGTAATAAATTGTACT- 76687

QY 922 GTATTGCGATTTCGAAATGATGTTGTTTACGCTCTTCCACAAAAAAGAAAA 981
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Db 76688 GTATTGCGATTTCGCAATTAGTAGATTCTAGCTGCTTCCACAGAAAGGAAAA 76747

QY 982 AAAAAA 991
| | | | |
Db 76748 AAATGGATAA 76757

RESULT 23
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LOCUS AC018788 146883 bp DNA linear PRI 22-NOV-2001
DEFINITION Homo sapiens chromosome , clone RP11-24E9, complete sequence.
ACCESSION AC018788
VERSION AC018788.10 GI:16117587
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 146883)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G., Castle,A.,
Choepel,Y., Collangelo,M., Collins,S., Collymore,A., Cooke,P.,
Dearellano,K., Dewar,K., Domino,M., Doyle,N., Feneator,J.,
Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Landers,I., Lehoczy,J., Levine,R., Liu,C., Liu,G., Locke,K.,
Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
Meldrum,J., Meneus,L., Morrow,J., Naylor,J., Norman,C.H.,

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O'Connor,T., O'Donnell,P., Olivar,T.M., Peterson,K., Pierre,N.,
Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D., Roy,A.,
Santos,R., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Tirrell,A.,
Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J., Zimmer,A.
and Zody,M.
Direct Submission
Submitted (19-DEC-1999) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 146883)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
Choepel,Y., Collangelo,M., Collins,S., Collymore,A., Cooke,A.,
Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
Ferreira,P., FitzHugh,W., Gage,D., Galagan,J., Gardyna,S.,
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Hagos,B., Heaford,A., Horton,L., Hulme,W., Iliev,I., Johnson,R.,
Jones,C., Kamat,A., Karatas,A., Kells,C., LaRocque,K.,
Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,
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McCarthy,M., McEwan,P., McKernan,K., McPheeters,R., Meldrum,J.,
Meneus,L., Mihova,T., Mlenga,V., Murphy,T., Naylor,J., Nguyen,C.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schuback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (22-NOV-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Oct 13, 2001 this sequence version replaced gi:16041478.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center / MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence.submissions@genome.wi.mit.edu
----- Project Information
Center project name: L4443
Center clone name: 24_E_9
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39819..39874
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Query Match 6.2%; Score 61.8; DB 9; Length 146883;
Best Local Similarity 57.5%; Pred. No. 0.00035;
Matches 111; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

QY 796 AGGAGTTGGGGGTTCCAGGATAGGAGTGGAGGTCAGAGGACGCAAGCAGCAGCA 855
Db 100174 AGGAGTGGAGGGGGGAGAAATGGGAGATGAGGTGAGGATACCAATA 100115

QY 856 TGTAGATGAACCGTCCAGAGAGCAAGCAGGAGAGCTGCAGGCGCATCAGGCTGCA 915
Db 100114 TGTAAATAACAACCTCTGGAGACCCCTATGTACACCATGAGGACTATGTTTAAGAT 100055

QY 916 CTGTCGTATTGGAGTTCATGCAAAATGAGTGTGTTTAGTCTCTTGCACAAAAA 975
Db 100054 TGTACTGTATTGGAGTTCTGCTAAATGAGTATATTTAGTCTCTTGCACAAAAA 99995

QY 976 AAAAAA 988
Db 99994 AAAAAA 99982

RESULT 24
AP429315 125020 bp DNA linear PRI 18-JAN-2002
LOCUS Homo sapiens junctophilin 3 (JPH3) gene, partial cds.
DEFINITION AP429315
ACCESSION AP429315
VERSION AP429315.1 GI:17646244
KEYWORDS
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
REFERENCE 1 (bases 1 to 125020)
AUTHORS Holmes,S.E., O'Hearn,E., Rosenblatt,A., Callahan,C., Hwang,H.S.,
Ingersoll-Ashworth,R.G., Fleisher,A., Stevanin,G., Brice,A.,
Potter,N.T., Ross,C.A. and Margolis,R.L.
TITLE A repeat expansion in the gene encoding junctophilin-3 is
associated with Huntington disease-like 2
JOURNAL Nat. Genet. 29 (4), 377-378 (2001)
MEDLINE 21583737
PUBMED 11694876
REFERENCE 2 (bases 1 to 125020)
AUTHORS Holmes,S.E., Ingersoll-Ashworth,R.G., Ross,C.A. and Margolis,R.L.

```


JOURNAL	Submitted (24-AUG-2002)	Whitehead Institute/MIT Center for Genome
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COMMENT

Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 11, 2000 this sequence version replaced g1:7280308.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research

Web site: <http://www-seq.wi.mit.edu>

Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information
 Center project name: L4198

Center clone name: 276.N.1

----- Summary Statistics

Sequencing vector: M13; M77815; 100% of reads

Assembly: Dye-terminator Big Dye; 100% of reads

Assembly program: Phrap; version 0.960731

Consensus quality: 148809 bases at least Q40

Consensus quality: 157187 bases at least Q30

Consensus quality: 160434 bases at least Q20

Insert size: 168000; agarose-fp

Insert size: 162462; sum-of-contigs

Quality coverage: 3.6 in Q20 bases; agarose-fp

Quality coverage: 3.7 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 24 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

```

1 1218: contig of 1218 bp in length
1219 1318: gap of 100 bp
1319 2383: contig of 1065 bp in length
2384 2483: gap of 100 bp
2484 3871: contig of 1388 bp in length
3872 3971: gap of 100 bp
3972 4836: contig of 865 bp in length
4837 4937: gap of 100 bp
4938 6262: contig of 1325 bp in length
6263 6361: gap of 100 bp
6362 8332: contig of 1971 bp in length
8333 8433: gap of 100 bp
8434 11370: contig of 2938 bp in length
11371 11470: gap of 100 bp
11471 16136: contig of 4666 bp in length
16137 16236: gap of 100 bp
16237 20360: contig of 4724 bp in length
20361 21060: gap of 100 bp
21061 25287: contig of 4227 bp in length
25288 25387: gap of 100 bp
25388 32039: contig of 6652 bp in length
32040 32139: gap of 100 bp
32140 38768: contig of 6629 bp in length
38769 38869: gap of 100 bp
38870 43784: contig of 4916 bp in length
43785 43884: gap of 100 bp
43885 50401: contig of 6517 bp in length
50402 50501: gap of 100 bp
50502 57082: contig of 6581 bp in length
57083 57182: gap of 100 bp
57183 63917: contig of 6735 bp in length
63918 64017: gap of 100 bp
64018 71454: contig of 7437 bp in length
71455 71554: gap of 100 bp
71555 79670: contig of 8116 bp in length
79671 79770: gap of 100 bp
79771 87413: contig of 7643 bp in length
87414 93773: gap of 100 bp
87514 93773: contig of 6260 bp in length
93774 93873: gap of 100 bp

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* 93874 103590: contig of 9717 bp in length
* 103591 103690: gap of 100 bp
* 103691 118309: contig of 14619 bp in length
* 118310 118409: gap of 100 bp
* 118410 136058: contig of 17649 bp in length
* 136059 136158: gap of 100 bp
* 136159 164762: contig of 28604 bp in length.

```

FEATURES

Source

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1..164762
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="9"
/map="9"

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/clone="RP11-276N1"
/clone_lib="RPC1-11 Human Male BAC"

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1..1218
/note="assembly_fragment"
1319..2383
/note="assembly_fragment"
2484..3871
/note="assembly_fragment"
3972..4836
/note="assembly_fragment
clone_end:SP6
vector_side:right"
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/note="assembly_fragment"
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/note="assembly_fragment"
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/note="assembly_fragment"
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/note="assembly_fragment"
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/note="assembly_fragment"
21061..25287
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25388..32039
/note="assembly_fragment"
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38869..43784
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/note="assembly_fragment"
50502..57082
/note="assembly_fragment"
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/note="assembly_fragment"
64018..71454
/note="assembly_fragment"
71555..79670
/note="assembly_fragment"
79771..87413
/note="assembly_fragment"
87514..93773
/note="assembly_fragment
clone_end:T7
vector_side:left"
93874..103590
/note="assembly_fragment"
103691..118309
/note="assembly_fragment"
118410..136058
/note="assembly_fragment"
136159..164762
/note="assembly_fragment"

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misc_feature
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```

BASE COUNT 47298 a 33699 c 32793 g 48671 t 2301 others
ORIGIN

```

```

Query Match          6.1%; Score 60.8; DB 2; Length 164762;
Best Local Similarity 64.1%; Fred. No. 0.0006;

```

Matches 109; Conservative 0; Mismatches 57; Indels 4; Gaps 1;

QY 814 GGATAGGGGAATGGGAGGTCAGAGGACGCAAGCAGCAGCCATGTAGATGAACCGTCCA 873

Db 64641 GAAATGGGAGATGTAGGTGTCAGAGGACACAAAGTAAACACATGTAGGTGAACAGTCT 64700

QY 874 GAGAGCCAAAGCAGCGGACAGGAGTCTAGGCGCATCAGCGTGCATGTTCTGATTTGGAGTT 933

Db 64701 AGAGATCAAAATGTACGTGAAGATTACAGATAATAAAATTGTACT---GTATTGTGATT 64756

QY 934 CATGCAAAATGAGTGTGTTTGTAGTCTCTTGGCCACAAAAAATAAAAA 983

Db 64757 CCTGCTAAATGAGTAGACTTTAGCTATTCTTGCCACAAAAAATAAAAA 64806

RESULT 26

AL353764/c

LOCUS

DEFINITION Human DNA sequence from clone RP11-440G5 on chromosome 9, complete sequence.

ACCESSION AL353764

VERSION AL353764.9 GI:14272263

KEYWORDS HTG.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE

AUTHORS Tracey,A.

TITLE Direct Submission

JOURNAL Submitted (01-JUN-2001) Sanger Centre, Hinxton, Cambridgeshire, CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk

COMMENT On May 31, 2001 this sequence version replaced gi:13990016. During sequence assembly data is compared from overlapping clones. Where differences are found these are annotated as variations together with a note of the overlapping clone name. Note that the variation annotation may not be found in the sequence submission corresponding to the overlapping clone, as we submit sequences with only a small overlap as described above.

This sequence was finished as follows unless otherwise noted: all regions were either double-stranded or sequenced with an alternate chemistry or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by at least one plasmid subclone or more than one M13 subclone, and the assembly was confirmed by restriction digest. The following abbreviations are used to associate primary accession numbers given in the feature table with their source databases: Em: EMBL; Sw: SWISSPROT; Tr: TrEMBL; Wp: WORMPEP; Information on the WORMPEP database can be found at

http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence was generated from part of bacterial clone contigs of human chromosome 9, constructed by the Sanger Centre Chromosome 9 Mapping Group. Further information can be found at

http://www.sanger.ac.uk/HGP/Chr9

RP11-440G5 is from the library RPCT-11.2 constructed by the group of Pieter de Jong. For further details see

http://www.chori.org/bacpac/home.htm

VECTOR: pBACes.6

This sequence is the entire insert of clone RP11-440G5 The true left end of clone RP11-818 is at 105684 in this sequence. The true right end of clone RP11-446A5 is at 70776 in this sequence.

FEATURES

Location/Qualifiers

1..170898

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="9"

/clone="RP11-440G5"

/clone_lib="RPCT-11.2"

BASE COUNT 48782 a 34836 c 35124 g 52156 t

ORIGIN

Query Match 6.1%; Score 60.8; DB 9; Length 170898;

Best Local Similarity 64.1%; Pred. No. 0.0006;

Matches 109; Conservative 0; Mismatches 57; Indels 4; Gaps 1;

QY 814 GGATAGGGGAATGGGAGGTCAGAGGACGCAAGCAGCAGCCATGTAGATGAACCGTCCA 873

Db 69053 GAAATGGGAGATGTAGGTGTCAGAGGACACAAAGTAAACACATGTAGGTGAACAGTCT 68994

QY 874 GAGAGCCAAAGCAGCGGACAGGAGTCTAGGCGCATCAGCGTGCATGTTCTGATTTGGAGTT 933

Db 68993 AGAGATCAAAATGTACGTGAAGATTACAGATAATAAAATTGTACT---GTATTGTGATT 68938

QY 934 CATGCAAAATGAGTGTGTTTGTAGTCTCTTGGCCACAAAAAATAAAAA 983

Db 68937 CCTGCTAAATGAGTAGACTTTAGCTATTCTTGCCACAAAAAATAAAAA 68888

RESULT 27

AL353764

LOCUS

DEFINITION Homo sapiens BAC clone RP11-1N7 from 2, complete sequence.

ACCESSION AC104684

VERSION AC104684.3 GI:18464246

KEYWORDS HTG.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE

AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

TITLE 1 (bases 1 to 129770)

JOURNAL Sulston,J.E. and Waterston,R.

MEDLINE Toward a complete human genome sequence

PUBMED Genome Res. 8 (11), 1097-1108 (1998)

REFERENCE

AUTHORS Belter,E., Meyer,R. and Creason,K.

TITLE The sequence of Homo sapiens BAC clone RP11-1N7

JOURNAL Unpublished (2001)

AUTHORS Waterston,R.H.

TITLE Direct Submission

JOURNAL Submitted (19-DEC-2001) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE

AUTHORS Waterston,R.H.

TITLE Direct Submission

JOURNAL Submitted (01-FEB-2002) Genome Sequencing Center, Washington University School of Medicine, 4444 Forest Park Parkway, St. Louis, MO 63108, USA

REFERENCE

AUTHORS Waterston,R.

TITLE Direct Submission

JOURNAL Submitted (21-FEB-2002) Department of Genetics, Washington University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA

COMMENT On Feb 1, 2002 this sequence version replaced gi:18250104.

Center: Washington University Genome Sequencing Center

Center code: WUGSC

Web site: http://genome.wustl.edu/gsc

Contact: sapiens@wustl.edu

Summary Statistics

Center project name: H_NH0001N07

NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate

chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/qsc>

SOURCE INFORMATION:

The RPI-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Moon, P. Y., Zhao, B., Eschgen, E., Tateno, M., Catanesi, J. and de Jong, P. J. (1998). An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at <http://www.chori.org>
VECTOR: DEACe3.6

VECTOR: pBACE3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is XXcos-2052C30; the clone sequenced to the right is RP11-356M6, 2000 bp overlap. Actual start of this clone is at base position 1 of RP11-IN7.

Sequence derived from one plasmid subclone, base position 94919 to 94957.

Data from AC067919 and AC097644 was used to finish this clone, AC104684.

Polymorphisms have been identified between AC067919, AC079779 and AC104684.

The sequence of AC021641 has been incorporated into AC104684.

FEATURES

location/Qualifiers	source
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/organism="Homo sapiens"	
/mol_type="genomic DNA"	
/db_xref="taxon:9606"	
/chromosome="2"	
/map="2"	
/clone="RP11-IN7"	
/clone_lib="RPC1-11"	
1..77	
/rpt_family="L1"	repeat_region
78..340	repeat_region
/rpt_family="Alu"	repeat_region
341..668	repeat_region
/rpt_family="L1"	repeat_region
661..932	repeat_region
/rpt_family="L1"	repeat_region
944..1216	repeat_region
/rpt_family="Alu"	repeat_region
1231..1277	repeat_region
/rpt_family="(TAAA)n"	repeat_region
1260..1341	repeat_region
/rpt_family="L1"	repeat_region
1342..1566	repeat_region
/rpt_family="Alu"	repeat_region
1567..1624	repeat_region
/rpt_family="L1"	repeat_region
1615..1636	repeat_region
/rpt_family="AT_rich"	repeat_region
1651..1778	repeat_region
/rpt_family="Alu"	repeat_region
3395..3458	repeat_region
/rpt_family="AchHobo"	repeat_region
3459..3749	repeat_region
/rpt_family="Alu"	repeat_region
3750..3801	repeat_region

repeat_region 29751..30738

Query Match 6.1%; Score 60.4; DB 9; Length 129770;
 Best Local Similarity 56.6%; Pred. No. 0.00076;
 Matches 112; Conservative 0; Mismatches 86; Indels 0; Gaps 0;
 QY 794 AGAGAGTTTGGGGTTCAGATAGGAATGGGAGGTACAGAGCCCAAGCAGCAGC 853
 Db 77435 AGGGGAGCTGGAGAGGGTGGGGGATGGGAAGATAGTGCAGAGGTACAAAGTAGCAGG 77494
 QY 854 CATGTAGATCAACCGTCCAGAGAGCCCAAGCAGCAGGACTGCAGGCCCATCAGCGTG 913
 Db 77495 TGTGTGGATTAACAGTCCAGAGATCTCATGTACAATGAGACATAGTGTAAAAA 77554
 QY 914 CACTGTTCTGATTTGGAGTTCATGCAAAATGAGTGTGTTTGTAGTGTCTTTGCCACAAA 973
 Db 77555 ATTGTGCTGATTTGGATTTCTGATAAATGAGTGTGATTGTAGTTGCTATTACCACAAA 77614
 QY 974 AAAAAAAAAAAAAAAAAA 991
 Db 77615 CAAAACAAAACAAAAA 77632

RESULT 28

AC067919/C
 LOCUS AC067919 171655 bp DNA linear HTG 26-MAY-2000
 DEFINITION Homo sapiens clone RP11-29216, WORKING DRAFT SEQUENCE, 20 unordered pieces.
 ACCESSION AC067919
 VERSION AC067919.2 GI:8076830
 KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 1 (bases 1 to 171655)
 Birren,B., Linton,L., Nusbaum,C. and Lander,E.
 Homo sapiens, clone RP11-29216
 Unpublished
 2 (bases 1 to 171655)
 Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N., Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F., Boguslavsky,L., Boukhalter,B., Brown,A., Burkett,G., Camporeale,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Domino,M., Doyle,M., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L., Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L., Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J., LaRoque,K., Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N., McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R., Meldrum,J., Meneus,L., Mihova,T., Miranda,C., Mlenga,V., Morrow,J., Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D., Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J., Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J., Zimmer,A. and Zody,M.

TITLE

JOURNAL Submitted (27-APR-2000) Whitehead Institute/MIT Center for Genome Research, 320 Charles Street, Cambridge, MA 02141, USA
 COMMENT On May 25, 2000 this sequence version replaced gi:7652358.
 All repeats were identified using RepeatMasker:

Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html

----- Genome Center

Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu

----- Project Information
 Center project name: I6333
 Center clone name: 292 I 6
 ----- Summary Statistics
 Sequencing vector: M13; M77815; 100% of reads
 Chemistry: Dye-terminator Big Dye; 100% of reads
 Assembly program: Phrap; version 0.960731
 Consensus quality: 161528 bases at least Q40
 Consensus quality: 167303 bases at least Q30
 Consensus quality: 168939 bases at least Q20
 Insert size: 173000; agarose-fp
 Insert size: 169755; sum-of-contigs
 Quality coverage: 4.1 in Q20 bases; agarose-fp
 Quality coverage: 4.1 in Q20 bases; sum-of-contigs

 * NOTE: This is a 'working draft' sequence. It currently
 * consists of 20 contigs. The true order of the pieces
 * is not known and their order in this sequence record is
 * arbitrary. Gaps between the contigs are represented as
 * runs of N, but the exact sizes of the gaps are unknown.
 * This record will be updated with the finished sequence
 * as soon as it is available and the accession number will
 * be preserved.

1 220: contig of 220 bp in length
 * 221 320: gap of 100 bp
 * 321 4001: contig of 3681 bp in length
 * 4002 4101: gap of 100 bp
 * 4102 7945: contig of 3844 bp in length
 * 7946 8045: gap of 100 bp
 * 8046 11450: contig of 3405 bp in length
 * 11451 11550: gap of 100 bp
 * 11551 16217: contig of 4567 bp in length
 * 16218 20480: contig of 4263 bp in length
 * 20481 20580: gap of 100 bp
 * 20581 25760: contig of 5180 bp in length
 * 25761 25860: gap of 100 bp
 * 25861 31039: contig of 5179 bp in length
 * 31040 31139: gap of 100 bp
 * 31140 35820: contig of 4681 bp in length
 * 35821 35920: gap of 100 bp
 * 35921 42490: contig of 6570 bp in length
 * 42491 42590: gap of 100 bp
 * 42591 49780: contig of 7190 bp in length
 * 49781 49880: gap of 100 bp
 * 49881 57796: contig of 7916 bp in length
 * 57797 57896: gap of 100 bp
 * 57897 67541: contig of 9645 bp in length
 * 67542 67641: gap of 100 bp
 * 67642 76045: contig of 8404 bp in length
 * 76046 76145: gap of 100 bp
 * 76146 85834: contig of 9689 bp in length
 * 85835 85934: gap of 100 bp
 * 85935 95701: contig of 9767 bp in length
 * 95702 95801: gap of 100 bp
 * 95802 107607: contig of 11806 bp in length
 * 107608 107707: gap of 100 bp
 * 107708 123352: contig of 15645 bp in length
 * 123353 123452: gap of 100 bp
 * 123453 141544: contig of 18092 bp in length
 * 141545 141644: gap of 100 bp
 * 141645 171655: contig of 30011 bp in length.

FEATURES

Source

1. 171655
 /organism="Homo sapiens"
 /mol_type="genomic DNA"
 /db_xref="taxon:9606"
 /clone="RP11-29216"
 /clone_lib="RPC1-11 Human Male BAC"
 /notes="assembly_fragment
 clone end:T7
 vector_side:left"

misc_feature

1. 220
 /notes="assembly_fragment
 clone end:T7
 vector_side:left"

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misc_feature      321..4001
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                  /note="assembly_fragment"
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                  /note="assembly_fragment"
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                  /note="assembly_fragment"
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                  /note="assembly_fragment"
misc_feature      20581..25760
                  /note="assembly_fragment"
misc_feature      25861..31039
                  /note="assembly_fragment"
misc_feature      31140..35820
                  /note="assembly_fragment"
                  clone_end:SP6
                  vector_side:right"
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misc_feature      57897..67541
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misc_feature      67642..76045
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misc_feature      95802..107607
                  /note="assembly_fragment"
misc_feature      107708..123352
                  /note="assembly_fragment"
misc_feature      123453..141544
                  /note="assembly_fragment"
misc_feature      141645..171655
                  /note="assembly_fragment"
BASE COUNT      50815 a 35321 c 34768 g 48847 t 1904 others
ORIGIN
Query Match      6.1%; Score 60.4; DB 2; Length 171655;
Best Local Similarity 56.6%; Pred. No. 0.00075;
Matches 112; Conservative 0; Mismatches 86; Indels 0; Gaps 0;
QY 794 AGAGGAGTTTGGGGGTTTCAGGATAGGGAATGGGAGGTCAGAGGACGCAAGACGACG 853
Db 150859 AGGGGACTGGAGAGGGTGGGGATGGGAGATATAGGTCAGAGTACAAAGTAGCAGG 150800
QY 854 CATGTAGATGACCGTCCAGAGAGCCAAAGCAGGAGGACTGCAGGCCATCAGCGTG 913
Db 150799 TGTGTGGGATAAACAAGTCCAGAGATCTCATGTACAACATGAGGACTAGTGTAAAAA 150740
QY 914 CACTGTTCTATTTCGAGTTCATGAAATGATGTTGTTTTCAGTCTCTTCCCAAAA 973
Db 150739 ATTGTGCTGTTATTGGAATCTCGTAATATGATGTTTATTGTTGCTATTACCAAAA 150680
QY 974 AAAAAAAAAAAAAAAAAA 991
Db 150679 CAACAACAAAAA 150662

RESULT 29
AC010072
LOCUS      AC010072      124347 bp      DNA      linear      PRI 19-NOV-1999
DEFINITION Homo sapiens chromosome 14q31 clone CTD-217314 containing TSHR
            gene, partial cds; and unknown gene, complete sequence.
ACCESSION  AC010072
VERSION    AC010072.5      GI:6453843
KEYWORDS   HTG.
SOURCE     Homo sapiens (human)

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ORGANISM

Homo sapiens

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.

REFERENCE

AUTHORS

1 (bases 1 to 124347)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B., Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G., James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T. and Hood,L.

TITLE

JOURNAL

Unpublished
Sequencing of human chromosome 14

REFERENCE

AUTHORS

2 (bases 1 to 124347)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B., Bloom,S., Dors,M., Dickhoff,R., Harrison,G., James,R., Lasky,S., Madan,A., Ratcliffe,A., Shaffer,T. and Hood,L.

TITLE

JOURNAL

Submitted (11-SEP-1999) Multimegabase Sequencing Center, University of Washington, PO BOX 357730, Seattle, WA 98195, USA

REFERENCE

AUTHORS

3 (bases 1 to 124347)
Rowen,L., Madan,A., Qin,S., Abbasi,N., Baradarani,L., Birditt,B., Bloom,S., Dors,M., Dickhoff,R., Fleetwood,P., Harrison,G., James,R., Kaur,A., Madan,A., Owen,M.P., Ratcliffe,A., Shaffer,T. and Hood,L.

TITLE

JOURNAL

Direct Submission
Submitted (19-NOV-1999) Multimegabase Sequencing Center, University of Washington, PO BOX 357730, Seattle, WA 98195, USA
On Nov 19, 1999 this sequence version replaced gi:6114900.

COMMENT

----- Genome Center

Center: Multimegabase Sequencing Center

Center code: UWNMC

Web site: http://chroma.mbt.washington.edu/msg_www

Contact: leerowen@u.washington.edu

----- Summary Statistics

Sequencing vector: pUC18; L08752

Chemistry: Big Dye terminators and primers

Assembly program: Phrap; version 0.990399

FEATURES

source

Location/Qualifiers

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/organism="Homo sapiens"

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/chromosome="14"

/map="14q31"

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/clone_lib="CalTech human BAC library D"

/note="This clone overlaps RP11-114N19, Accession

AC007262"

complement(join(42575..42673,43714..43786,47934..48025,

52809..52927,53951..54077,62318..62371))

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A1554064, the closest BLASTX similarity is to a

drosohila potassium channel protein"

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95810..95860

/note="Low quality data"

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/product="thyroid stimulating hormone receptor"

/note="This is the 5' end of TSHR, found in M73745"

103677..103846

/note="thyroid stimulating hormone receptor"

/codon_start=1

/product="TSHR"

/protein_id="AAF09032.1"

/db_xref="GI:6453844"

/translation="MRPADLLQLVLLLDLPRLDGMGCGSPPCFCEHQEDFRVTKOI

GRIFSLPPSTQTL"

unsure

mRNA

CDS

estimates computed by the Phrap assembly program.

All manually edited bases have been reduced to quality zero.

Quality levels above 40 are expected to have less than

1 error in 10,000 bp.

Base-by-base quality values are not generally visible from the

GenBank flat file format but are available as part

of this entry's ASN.1 file.

This sequence was finished as follows unless otherwise noted:
all regions were either double-stranded or sequenced with an
alternate chemistry or covered by high quality data (i.e., Phred
quality >= 30); an attempt was made to resolve all sequencing
problems, such as compressions and repeats; all regions were
covered by at least one plasmid subclone or more than one M13
subclone; and the assembly was confirmed by restriction digest.

Sequence Validation:

This sequence has been validated by Multiple Complete Digest
fingerprinting. Comparison of the experimentally derived digest
fragments with sequence-predicted fragments is given below.
The electronically-digested sequence consists of both insert and
vector, in order to accurately represent the entire circular BAC.
Small fragments below a variable cutoff (approximately 400-800 bp)
are not resolved in the fingerprint and hence do not appear
in the table. There are no significant remaining discrepancies
between the experimental and predicted values. Uniquely ordered
fragments are separated by dashed lines.

BcoRI

HindIII

BglII

SeqDerMap	FngPrnt	SeqDerMap	FngPrnt	SeqDerMap	FngPrnt
8696	8949	4503	4596	6422	6405
6	<800	6382	6584	2067	2089
6912	6827	512	<800	6249	6405
2142	2138	449	<800	470	<800
1633	1601	2282	2301	3028	2892
185	<800	404	<800	10089	9902
33	<800	3748	3800	2765	2775
235	<800	882	868	1438	1439
986	959	496	<800	5480	5750
5804	5786	1453	1429	8688	8596
166	<800	207	<800	2063	2089
2553	2584	4653	4596	7872	7784
73	<800	5302	5418	2057	2089
8410	8322	2615	2732	1970	2089
1588	1601	423	<800	344	<800
1392	1382	156	<800	5861	5750
3402	3455	3436	3385	965	988
147	<800	601	<800	4108	3983
1254	1250	52	<800	3172	3167
973	959	2019	2022	1145	1094

161	<800	3795	3800	1593	1595
313	<800	1540	1503	5146	5297
3713	3718	10075	9954	3644	3587
2627	2584	5447	5418	2365	2381
2804	2824	3913	3800	6608	6405
782	806	2683	2867	3181	3167
305	<800	622	<800	1074	988
2256	2264	1811	1749	1702	1736
4178	4181	1519	1503	759	<800
964	959	2068	2022	9566	9902
1486	1472	3072	3098	1419	1439
3586	3455	84	<800	9853	9902
6519	6434	381	<800	783	788
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1241	1250	2365	2540	2875	2892
3064	3099	5565	5418	205	<800
3415	3455	5400	5418	1223	1144
4194	4181	8055	8069	2821	2892
2586	2584	2087	2022	238	<800
2823	2824	857	868	939	988
3419	3455	613	<800	3195	3167
1501	1472	837	868	5065	4986
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4338	4390	6244	6292	5864	5750
5735	5786	3337	3385	1092	1094
4197	4181	2253	2301	379	<800
10513	10499	2916	2867	5118	4986
907	959	277	<800	989	988
842	806	1915	1912		
905	959	541	<800		
78	<800	574	<800		
12446	12330	7293	7311		
1636	1601	9275	9237		
1342	1382	2546	2622		
2807	2824	806	868		


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123684 atggctgcmg(n)ngmggcggg
123685 tggctgcmg(n)ngmggcggg
123686 gctgcmg(n)ngmggcggg
123687 gctgcmg(n)ngmggcggg
123688 ctgcmg(n)ngmggcggg
123702 ggggtgcmg(n)ngmggcggg
123720 ccttctctc(g)ctgcmg
123731 ctgcmg(n)ngmggcggg
123763 ctgcmg(n)ngmggcggg
123763 taggaagga(n)tggtctggac
123763 aaaaaaaa(n)cgagaagaa
123764 aaaaaaaa(n)cgagaagaa
123764 ggggtcagga(n)ttcgagaca
123764 ttctcaga(n)atcattct
123764 aaaaaaaa(n)aaagttaagc
123764 tactcataa(n)taattaccac
123764 taggaagga(n)tggtctggac
123764 agctgaaga(n)tcagctct
123764 ggaattct(n)acnccatc
123764 attcctact(n)ccatccagg
123764 ctttccca(n)gtcatttaag
123764 cacttatga(n)gnaataaam
123764 ccttatgag(n)gaataaung
123764 angnaataa(n)naaggcagc
123764 ngnaataa(n)ggggcagc
123764 acaacagca(n)actgctggt
123764 actgtgga(n)actggcgaag
123764 aagcctgtg(n)ttcagacgt

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----- Distribution of Quality < 40 Bases -----

# bases	5	10	15	20	25	30	35	40
1000								
900								
800								
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600								
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100								
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Version: 1.01 qxfo.
Location/Qualifiers
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/organism="Homo sapiens"
/mol_type="genomic DNA"
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/chromosome="3"
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complement(40. .496)
/rpt_family="L1M4"
1423. .1452
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1705. .2095
/rpt_family="L1R16A"
complement(2399. .2430)
/rpt_family="L2"
complement(2431. .2954)
/rpt_family="MLT1F"
complement(2955. .3056)
/rpt_family="L2"
complement(3109. .3199)
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5566. .5781
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6399. .6418

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FEATURES
source

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repeat_region
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repeat_region
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repeat_region

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Query Match 5.9%; Score 58.6; DB 9; Length 194635;
Best Local Similarity 61.0%; Pred. No. 0.002;
Matches 114; Conservative 0; Mismatches 69; Indels 4; Gaps 1;

797 GGAGTTTGGGGGTCAGGATAGGAATGGGAGGTCAGAGGAGCGGAAAGCAGCAGCCAT 856
54877 GGATGTGGGAGGCGGAGGAATGAGGATGTTAGGTCAGAGATACAAATGGCAATAT 54936
857 GTAGATGAACCTCCAGAGAGCAACGCGGAGGAGGATGCGAGGCGCATCAGCGTGCAC 916
54937 GTATGATTACAGGTGAGTCTTAATACAAATGAGAAATTATAGGTAATAAATTTGTAC 54996
917 TGTTCTGTTTGGGATTCATGCAAAATGAGTGTGTTTAGTCTCTTGGCCACAAAAA 976
54997 T-----GTATGTGAGATTCATGCTAAATGAGTAGATTTACCTCTTGGCCACAAAAA 55052
977 AAAAAA 983
55053 AACAGAA 55059

RESULT 32
HSM130AC2/c

LOCUS HSM130AC2 4950 bp mRNA linear PRI 22-NOV-1996
DEFINITION H.sapiens mRNA for M130 antigen cytoplasmic variant 2.
ACCESSION Z22970
VERSION Z22970.1 GI:312145
KEYWORDS antigen; antigen M130.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 4950)
AUTHORS Law S.K., Micklem, K.J., Shaw, J.M., Zhang, X.P., Dong, Y., Willis, A.C. and Mason, D.Y.
TITLE A new macrophage differentiation antigen which is a member of the scavenger receptor superfamily
Eur. J. Immunol. 23 (9), 2320-2325 (1993)
JOURNAL 93380506
MEDLINE 8370408
REFERENCE 2 (bases 1 to 4950)
AUTHORS Micklem, K.K.
TITLE Direct Submission
JOURNAL Submitted (14-JUN-1993) Kingsley K.J. Micklem, Nuffield Department of Pathology, University of Oxford, Level 1, Maternity Block, John Radcliffe, Hospital, Headington, Oxford, OX3 9DU, United Kingdom
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102. .3572
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DDLICNGNESALMNCKHQGWKHNCKDAEDAGVTCRSGADLSRLDVGTVCSGRLEV
RFQGEWGTICDDGWDSDYAAVACQKQCPATAVTAIGRVNASKGFHGLWLDVSQGH
PAWQCKHEWKGHYCNHEDAGVTCDSGLRLRGGSRCACTVEVELLLGKV
CDRGWGLKADVVCRLQCGSALTKSYQVYSKIQATNTWLFSLSCNGNETSLWCKNW
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CDS

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102..221
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/evidence=experimental
polyA_signal 4924..4929
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polyA_signal 4932..4937
polyA_site 4947
BASE COUNT 1348 a 970 c 1277 g 1355 t
ORIGIN
Query Match 5.9%; Score 58.4; DB 9; Length 4950;
Best Local Similarity 59.0%; Pred. No. 0.0025;
Matches 118; Conservative 0; Mismatches 81; Indels 1; Gaps 1;
QY 787 TGAGTGAGAGGAGTTGGGGGTTTCAGATAGGAATGGGAGGTCTAGAGACCAAG 846
Db 4080 TGACAGGAATGGGAATGAAGAACTGTGAAGACGTATGTAGGTCTAGAGATACAAG 4021
QY 847 CAGCAGCCATGTAGAAATGAACCTGCAGAGACCAAGCAGCAGGACTGCAGGCCAT 906
Db 4020 TAGCAGATACGTAGATGAACAGTCTAGAGTCTTAATGTACAAATGAGATATAGGT 3961
QY 907 CAGCGTGCACTTTCGTATTGGAGTTTCATGCAMAAATGAGTGTGTTTGTGCTCTTTC 966
Db 3960 AATAACATTGTGCT-GTATGTAGGATTCATGCTAAGTGTAGTGTGTTTGTGCTCTTTC 3902
QY 967 CACAAAAAATAAAAAA 986
Db 3901 CACAGAACAACAAAAA 3882
RESULT 33
AC131206 177260 bp DNA linear PRI 31-DEC-2002
LOCUS Homo sapiens 12 BAC RP11-157G21 (Roswell Park Cancer Institute
DEFINITION Human BAC Library) complete sequence.
AC131206
VERSION AC131206.2 GI:22549585
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 177260)
Muzny D.M., Adams C., Adio-Oduola B., Ali-oshman, P.R., Allen, C.,
Albrooks, S.L., Amarantunge, H.C., Are, J.R., Ayale, M., Banks, T.,
Barbaria, J., Benton, J., Bimaga, K., Blankenburg, K., Bomlin, D.,
Bouck, J., Bowie, S., Brileva, M., Brown, E., Brown, M., Bryant, N.P.,
Buhay, C., Burch, P., Burkett, C., Burrell, K.L., Byrd, N.C.,
Carroll, T.F., Carter, M., Cavazos, S.R., Chacko, J., Chavez, D.,
Chen, G., Chen, R., Chen, Z., Chiu, D., Chowdhry, I., Christopoulos, C.,
Cleveland, C.D., Cox, C., Coyle, M.D., Dathorne, S.R., David, R.,
Davila, M.L., Davis, C., Davy-Carroll, L., Dederich, D.A.,
Dolaney, K.R., Delgado, O., Denn, A.L., Ding, Y., Dinh, H.H.,
Douthwaite, K.J., Draper, H., Dugan-Rocha, S., Durbin, K.J.,
Earnhart, C., Edgar, D., Edwards, C.C., Elhaj, C., Emerling, S.,
Escotto, M., Falla, T., Ferraguto, D., Flagg, N., Ford, J., Foster, P.,
Frantz, P., Gabisi, A., Gao, J., Garcia, A., Garner, T., Garza, N.,
Gill, R., Gorrell, J.H., Guevara, W., Gunaratne, P., Hale, S.,
Hamilton, K., Han, J., Harris, C., Harris, K., Hart, M., Haviak, P.,

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Hawes, A., Hernandez, J., Hernandez, O., Hodgson, A., Hoques, M., Holloway, C., Hollins, B., Honsi, F., Howard, S., Huber, J., Hulyk, S., Hume, J., Ioshikhes, I., Jackson, L.E., Jacobson, B., Jia, Y., Johnson, R., Jolivet, S., Joudah, S., Karlson, E., Kelly, S., Khan, U., King, L., Korvah, J., Kovar, C., Kratovic, J., Kureshi, A., Landry, N., Leal, B., Lee, B., Lewis, L., Lewis, L., Li, J., Li, Z., Lichtarge, O., Lieu, C., Liu, J., Liu, W., Loulseghe, H., Lozada, R., Lu, X., Lucier, A., Lucier, R., Luna, R., Ma, J., Maheshwari, M., Mapua, P., Marandel, I., Martin, R., Martindale, A., Martinez, E., Massey, E., Mawhney, E., McLeod, M.P., Meador, M., Mei, G., Merscher, S., Merzhat, K., Montgomery, K.T., Morgan, M., Morris, S., Moser, M., Mohabbat, K., Nelson, D., Newton, J., Newton, N., Nguyen, A., Nguyen, N., Nguyen, N., Nickerson, E., Nwokenkwo, S., Ogih, M., Okwuonu, G., Oragunye, N., Oviado, R., Pace, A., Payton, B., Peery, J., Perez, L., Peters, L., Pickens, R., Primus, E., Pu, L.L., Quiles, M., Ren, Y., Rives, M., Rojas, A., Rojibokan, I., Rolfe, M., Ruiz, S., Savory, G., Scherzer, S., Scott, G., Shen, H., Shim, C., Shoostari, N., Sisson, I., Sodergren, E., Sonaika, T., Sparks, A., Stanley, H., Stone, H., Sutton, A., Svatek, A., Tabor, P., Tamerisa, A., Tamerisa, K., Tang, H., Tansey, J., Taylor, C., Taylor, T., Telford, B., Thomas, N., Thomas, S., Usmani, K., Vasquez, L., Vera, V., Villalon, D., Vinson, R., Wang, Q., Wang, S., Ward-Moore, S., Warren, R., Washington, C., Watlington, S., Williams, G., Williamson, A., Wleczky, R., Wooden, S., Worley, K., Wu, C., Wu, Y., Wu, Y.F., Zhou, J., Zorrilla, S., Zucherlapati, R., Weinstock, G. and Gibbs, R.

Direct Submission

Unpublished

2 (bases 1 to 177260)

Worley, K.C.

Direct Submission

Submitted (18-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

3 (bases 1 to 177260)

Worley, K.C.

Direct Submission

Submitted (30-AUG-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

4 (bases 1 to 177260)

Worley, K.C.

Direct Submission

Submitted (25-SEP-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

5 (bases 1 to 177260)

Worley, K.C.

Direct Submission

Submitted (31-DEC-2002) Human Genome Sequencing Center, Department of Molecular and Human Genetics, Baylor College of Medicine, One Baylor Plaza, Houston, TX 77030, USA

On Aug 30, 2002 this sequence version replaced gi:22296911.

INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email

gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSs are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence

continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL: <http://www.hgsc.bcm.tmc.edu:8088/quality.info/genbank.annotation.ht> ml.

FEATURES

Source

Location/Qualifiers

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/function="clone overlap"

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/note="overlaps bases 1..2025 of clone AC006927"

/function="clone overlap"

BASE COUNT 59476 a 33910 c 33289 g 50585 t

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Query Match 5.9%; Score 58.4; DB 9; Length 177260;

Best Local Similarity 59.0%; Pred. No. 0.0023;

Matches 118; Conservative 0; Mismatches 81; Indels 1; Gaps 1;

QY 787 TGAGTGAGGAGGAGTTGGGGGTTTCAGGATAGGGAATGGGAGGTTCAGAGCAGCAAG 846

Db 64219 TGACAGGAATGGGAATGGAAGAACTGTGAAGACGTATGTAGGTTCAGAGGATCAAG 64278

QY 847 CAGAGCCATGTAGATGAACCGTCCAGAGAGCCAGCAGCAGGAGCACTGCAGGCCAT 906

Db 64279 TAGCAGATACGTAGATGAACAGCTAGAGGTCTAATCTACACATGAGGATTATAGT 64338

QY 907 CAGCGTGACTTGTGTTATTCGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTTCG 966

Db 64339 AATAACATTGCTCT-GTATGTAGGATTCATGCTAAGTAGTAGTGTGTTTAGCTGCTTTCG 64337

QY 967 CACAAAAAATAAAAAA 986

Db 64398 CACAGACACACACAAAAA 64417

RESULT 34

AC023003
LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
REFERENCE
AUTHORS
TITLE
JOURNAL
REFERENCE
AUTHORS

AC023003
Homo sapiens clone RP11-270M20, WORKING DRAFT SEQUENCE, 12
unordered pieces.
AC023003
2 GI:7139812
HTG; HTGS_PHASE1; HTGS_DRAFT.
Homo sapiens (human)
Homo sapiens
Bukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 140055)
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Homo sapiens, clone RP11-270M20
Unpublished
2 (bases 1 to 140055)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Beckerly,R., Beda,F., Castle,A.,
Boguslavsky,L., Bouckgater,B., Brown,A., Burkett,G., Castle,A.,
Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
DeArelano,K., Dewar,K., Domino,M., Doyle,M., Fenestor,J.,
Ferreira,P., FitzHugh,W., Forrest,C., Gage,D., Galagan,J.,
Gardyna,S., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Johnson,R., Jones,C., Kann,L., Karatas,A., Klein,J.,
Lander,T., Lechoczy,J., Levine,R., Lieu,C., Liu,G., Locke,K.,
Macdonald,P., Marquis,N., McEwan,P., McGurk,A., McKernan,K.,
McPheeters,R., Meldrum,J., Meneus,L., Morrow,J., Naylor,J.,
Norman,C.H., O'Connor,T., O'Donnell,P., Olivari,T.M., Peterson,K.,
Pierre,N., Pisani,C., Pollara,V., Raymond,C., Riley,R., Rothman,D.,
Roy,A., Santos,R., Severy,P., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Vassiliev,H., Viel,R., Vo,A., Wu,X., Wyman,D., Ye,W.J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (07-FEB-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141 USA
On Mar 1, 2000 this sequence version replaced gi:6921586.
All repeats were identified using RepeatMasker:
Smit, A.F.A. & Green, P. (1996-1997)
<http://ftp.genome.washington.edu/RM/RepeatMasker.html>

Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIRK
Web site: <http://www-seq.wi.mit.edu>
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: 270 M 20
Center clone name: 16597
----- Summary Statistics

Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 133511 bases at least Q40
Consensus quality: 136178 bases at least Q30
Consensus quality: 137450 bases at least Q20
Insert size: 135000; agarose-fp
Insert size: 138955; sum-of-contigs
Quality coverage: 5.1 in Q20 bases; agarose-fp
Quality coverage: 5.1 in Q20 bases; sum-of-contigs

* NOTE: This is a 'working draft' sequence. It currently
* consists of 12 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 1060: contig of 1060 bp in length
* 1061 1160: gap of 100 bp
* 1161 3767: contig of 2607 bp in length
* 3768 3867: gap of 100 bp
* 3868 8490: contig of 4623 bp in length

*	8491	8590: gap of 100 bp	
*	8591	14297: contig of 5707 bp in length	
*	14298	14397: gap of 100 bp	
*	14398	20798: contig of 6401 bp in length	
*	20799	20898: gap of 100 bp	
*	20899	31912: contig of 11014 bp in length	
*	31913	32012: gap of 100 bp	
*	32013	43544: contig of 11532 bp in length	
*	43545	43644: gap of 100 bp	
*	43645	52029: contig of 8385 bp in length	
*	52030	52129: gap of 100 bp	
*	52130	67985: contig of 15856 bp in length	
*	67986	68085: gap of 100 bp	
*	68086	84647: contig of 18562 bp in length	
*	84648	84747: gap of 100 bp	
*	84748	108247: contig of 23500 bp in length	
*	108248	108347: gap of 100 bp	
*	108348	140055: contig of 31708 bp in length.	
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	/mol_type="Genomic DNA"		
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	/clone_lib="RPC1-11 Human Male BAC"		
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	/note="assembly_fragment"		
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	/note="assembly_fragment"		
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misc_feature	vector_side:left		
	52130. 67985		
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misc_feature	68086. 84647		
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misc_feature	84748. 108247		
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Best Local Similarity	61.9%;	Pred. No. 0.0029;	
Matches 125;	Conservative	0;	Mismatches 75; Indels 2; Gaps 2;
QY	787	TGAGTGAAGAGGAGTTGGGGGTTCCAGGATAGGGAATGGGAGGTTCAGAGACGCAAG	846
Db	15860	TGACAGAGGAGGTAGAAATGTCATAGGAATGGGAATATATAGGTCAGAGGACACAAAG	15919
QY	847	CACAGGCCATGTGAGATGAACC-GTCCAGAGAGCCAAAGCACGCGCAGGACTGCAGGCCA	905
Db	15920	CACAGGCCATGCAGGAGGACCAGAGTCTAGAGATCTTAAGCTACATTATGAGGACCATACAG	15979
QY	906	TCACGCTGCACCTTCGTATTGGAGTTTCATCAAAATGAGTGTGTTTACGTCGCTCTTG	965
Db	15980	GTAATTAAGACGGTACTGTGTATGGGATTTCATGCTAACTAGTA-GATTTGGCTGCTCTTG	16038
QY	966	CCACAAAAAAAAAAAAAAAAAA	987

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1745. .2410
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2411. .2705
/note="AluSg repeat: matches 1. .296 of consensus"
2706. .3163
/note="L1MC3 repeat: matches 7281. .7736 of consensus"
3183. .3479
/note="AluSg repeat: matches 1. .294 of consensus"
complement (4350. .4820)
/note="match: GSS: Em:AQ827082"
4999. .5225
/note="L2 repeat: matches 1328. .1564 of consensus"
5811. .6105
/note="AluSg repeat: matches 1. .295 of consensus"
6180. .6371
/note="L1MC3 repeat: matches 5725. .5926 of consensus"
6386. .6421
/note="18 copies 2 mer gt 97% conserved"
6673. .7475
/note="L2 repeat: matches 1777. .2614 of consensus"
7485. .7709
/note="MIR repeat: matches 21. .262 of consensus"
7733. .7783
/note="L2 repeat: matches 2648. .2705 of consensus"
8206. .8517
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8833. .10173
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10388. .10664
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11826. .12145
/note="AluSx repeat: matches 1. .294 of consensus"
12656. .12756
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12776. .13081
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complement (13145. .13540)
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13457. .13718
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14443. .14562
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14515. .14558
/note="22 copies 2 mer ag 77% conserved"
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80037. .80147, 88401. .88572, 97782. .97814, 97914. .98067,
107109. .107254))
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Tr:Q9W506 Tr:O61825 Sw:FL18490"
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/evidence=not_experimental
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/db_xref="GI:18070856"
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QDPDPSFIHNGHIIITYRPTVFCVLLGLLDGTAKARHPFVYVYGLKLFSPVFL
QSRDYLIVLFCFPAISLGLFPQINTFTVLLEIOIDMLFFGGSAVSITSAVSV
RSVLAALHVCFAKPEWPMOHIPALFSAFCGLVALSVHLRQSDPSVLSF
OCLRPKFLHONLAESAADPLPKMKDSYDVLKVDLLVCVAVLSFAVSAVSFVLS
LRPFLSIVLFALAGVGFVTHVYLPQLRKHHPWMLISHELKKNKTHEQREVDVAHLM

WFERLYVLOCFERKYLILPALILNALTIDAFILSNHRLGTHWDIFLMIAGMKLLRT
SECPYQPINLSFTVIFDFDYKDISSEFLDFMVSILFSK"
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16957. .17561
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20315. .20425
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21268. .21570
/note="AluSx repeat: matches 1. .297 of consensus"
21571. .21690
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22427. .23076
/note="L2 repeat: matches 1148. .1840 of consensus"
23241. .23347
/note="MER5A repeat: matches 4. .112 of consensus"
23351. .23520
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23676. .23858
/note="L2 repeat: matches 953. .1144 of consensus"
23868. .24080
/note="AluJb repeat: matches 87. .298 of consensus"
24270. .24836
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24837. .25148
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complement (24980. .25609)
/genes="dJ1016N21.1"
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/note="LIME repeat: matches 4886. .5093 of consensus"
25566. .25747
/note="MER97b repeat: matches 1. .187 of consensus"
complement (25605. .26037)
/genes="dJ1016N21.1"
/note="match: GSS: Em:AQ683895"
25810. .25859
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29471. .29912
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29498. .29966
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Query Match 5.8%; Score 58; DB 9; Length 151553;
Best Local Similarity 59.7%; Pred. No. 0.0029;
Matches 117; Conservative 0; Mismatches 75; Indels 4; Gaps 1;
QY 789 ACTGAAGAGGAGTTTGGGGGTTTCAGGATAGGAATGGGGAGTTCAGAGGACGCAAGCA 848
DB 47275 AGTTAACTATGGCATGGGGAGGTAGACATGGGGATATGTGTGTAGAAATTTATAAGTA 47334
QY 849 GCAGCCATGTAAGTAACCGTCCAGAGACCAACGACGACGATGCGAGCCATCA 908
DB 47335 GCAGATATGTAGGATAAAACAACCTCCAGCGATCTAATGTACATGAGAACTATAATTA 47394

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QY 909 GCGTCACTTGTGTTATTTGGATTCATGCAAAATGAGTGTGTTAGTCTTGCCCA 968
Db 47395 AACTGTACT-----ATATTGGGATTCCTGCAAAATAGTAGATTTAGTCTTGCCCA 47450

QY 969 CAAAAAATAAAAAA 984
Db 47451 CAAAAAATAAAAAA 47466

RESULT 36
AC019159/c
LOCUS AC019159 163085 bp DNA linear PRI 07-NOV-2001
DEFINITION Homo sapiens BAC clone RP11-56018 from 2, complete sequence.
ACCESSION AC019159
VERSION AC019159.8 GI:13677116
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Sulston,J.E. and Waterston,R.
Toward a complete human genome sequence
Genome Res. 8 (11), 1097-1108 (1998)
99063732
9847074
PUBMED
2 (bases 1 to 163085)
Goyea,E., Cotton,M., Spalding,L. and Lehnert,L.
The sequence of Homo sapiens BAC clone RP11-56018
Unpublished
3 (bases 1 to 163085)
Waterston,R.H.
Direct Submission
Submitted (30-DEC-1999) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
4 (bases 1 to 163085)
Waterston,R.H.
Direct Submission
Submitted (19-APR-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
5 (bases 1 to 163085)
Waterston,R.H.
Direct Submission
Submitted (20-APR-2001) Genome Sequencing Center, Washington
University School of Medicine, 4444 Forest Park Parkway, St. Louis,
MO 63108, USA
6 (bases 1 to 163085)
Waterston,R.
Direct Submission
Submitted (07-NOV-2001) Department of Genetics, Washington
University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
On Apr 19, 2001 this sequence version replaced gi:11276269.
----- Genome Center
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
----- Summary Statistics
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Center project name: H_NH0056018

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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence

from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RP11-11 human BAC library was made from the blood of one male donor, as described by Osoegawa, K., Woon, P. Y., Zhao, B., Frengen, E., Tatenio, M., Catanese, J. J., and de Jong, P. J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. Genomics 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is AC027111; the clone sequenced to the right is AC023003. Actual start of this clone is at base position 1 of RP11-56018; actual end is at base position 163085 of RP11-56018.

The sequence from base position 822245 to 82358 is derived from a single plasmid subclone. Size information based on PCR supports the assembly.

FEATURES

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	197. 373
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	8077. 8445
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Query Match 5.8%; Score 58; DB 9; Length 163085;
Best Local Similarity 61.9%; Pred. No. 0.0029;
Matches 125; Conservative 0; Mismatches 75; Indels 2; Gaps 2;

QY 787 TGAGTGAAGAGGAGTTGGGGGTTTCAGGATAGGGAATGGGAGGTTCAGAGGACGCAAG 846
Db 76533 TCACAGAGAGGGTAGAAATGCATAGGAATGGGAATATATAGGTTCAGAGGACCAAG 76474

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QY 847 CAGCAGCCATGTAGATGAACC-GTCCAGAGACCCAGACGCGCAGGAGCTGCAGGCCA 905
Db 76473 CAGCAGCCATGTAGAGGAGGACCAAGTCTAGAGATCTAAGGTACATTAAGGACCATACAG 76414
QY 906 TCAGCGTGCACCTGTTGCTATTGAGTTTCATGCMAAATCAGTGTGTGTTAGTGTGCTCTTG 965
Db 76413 GTAAATPAAGAGCGTACTGTGTGTATGGGATTCATGCTAAACTAGTA-GATTGGCTGCTCTTG 76355
QY 966 CCACAAAAAATAAAAAAAAAA 987
Db 76354 CCACAAAAAATAAAAAAAAAA 76333

RESULT 37
AC091178 99268 bp DNA linear PRI 01-APR-2002
LOCUS Homo sapiens chromosome 17, clone CTD-2206N4, complete sequence.
DEFINITION AC091178
ACCESSION AC091178
VERSION AC091178.7 GI:19852133
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Birren,B., Linton,L., Nusbaum,C. and Lander,E.
Unpublished
REFERENCE 1 (bases 1 to 99268)
Homo sapiens chromosome 17, clone CTD-2206N4
2 (bases 1 to 99268)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Boguslavskiy,L., Bouckgalter,B., Brown,A.,
Camarata,J., Campopiano,A., Chang,J., Choepel,Y., Colangelo,M.,
Collins,S., Collymore,A., Cooke,P., DeArellano,K., Dewar,K.,
Diaz,J.S., Dodge,S., Faro,S., Ferreira,P., Fitzhugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Hagos,B., Heaford,A., Horton,L., Hulme,W.,
Iliev,I., Johnson,R., Jones,C., Karatas,A., LaRocque,K.,
Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Liu,G.,
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Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Norbu,C.,
Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J.,
Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R., Seaman,S.,
Severy,P., Sougnez,C., Spencer,B., Stange-Thomann,N.,
Stojanovic,N., Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S.,
Theodore,J., Travers,M., Travis,N., Trigglio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (03-APR-2001) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 99268)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Ali,A., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Bloom,T., Boguslavskiy,L.,
Bouckgalter,B., Brown,A., Camarata,J., Campopiano,A., Chang,J.,
Chazaro,B., Choepel,Y., Colangelo,M., Collins,S., Collymore,A.,
Cooke,P., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
Kanat,A., Karatas,A., Kells,C., LaRocque,K., Lamazares,R.,
Landers,T., Lehoczy,J., Levine,R., Lindblad-Toh,K., Liu,G.,
Maclean,C., Macdonald,P., Major,J., Marquis,N., Matthews,C.,
McCarthy,M., McEwan,P., McKernan,K., Meldrim,J., Menes,L.,
Mihova,T., Mienga,V., Murphy,T., Naylor,J., Nguyen,C., Nicol,R.,
Norbu,C., Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D.,
Oliver,J., Peterson,K., Phunkhang,P., Pierre,N., Pollara,V.,
Raymond,C., Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P.,
Roman,J., Rosetti,M., Roy,A., Santos,R., Schauer,S., Schupback,R.,
Seaman,S., Severy,P., Spencer,B., Stange-Thomann,N., Stojanovic,N.,

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Strauss, N., Subramanian, A., Talamas, J., Tesfaye, S., Theodore, J.,
 Totham, K., Travers, M., Travis, N., Trigilio, J., Vassiliev, H.,
 Viel, R., Vo, A., Wilson, B., Wu, X., Wyman, D., Ye, W.-J., Young, G.,
 Zainoun, J., Zembek, L., Zimmer, A. and Zody, M.
 Direct Submission
 Submitted (01-APR-2002) Whitehead Institute/MIT Center for Genome
 Research, 320 Charles Street, Cambridge, MA 02141, USA
 On Apr 1, 2002 this sequence version replaced gi:18643501.
 All repeats were identified using RepeatMasker:
 Smit, A.F.A. & Green, P. (1996-1997)
 http://ftp.genome.washington.edu/RM/RepeatMasker.html
 ----- Genome Center
 Center: Whitehead Institute/ MIT Center for Genome Research
 Center code: WIBR
 Web site: http://www-seq.wi.mit.edu
 Contact: sequence_submissions@genome.wi.mit.edu
 ----- Project Information
 Center project name: L13055
 Center clone name: 2206_N_4

Only the first 99.2 kilobases of this clone are being submitted.
 The remainder of the clone is overlapped by either accession number
 AC006441 [WICGR project L515] or
 accession number AC004408 [WICGR project L309].

FEATURES

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1. 99268
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 /db_xref="taxon:9606"
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 /map="17"
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 1325..1501
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 /rpt_family="AluSg"
 1805..2245
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 2246..2567
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 2708..2999
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 complement(3903..3908)
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 complement(3915..3919)
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 6814..6824
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 7064..7069
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 7161..7227
 /rpt_family="(GGAA)n"
 7385..7474
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 complement(7510..7804)
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 7540..7551
 /note="<30 qual SNGL region"
 9096..9436
 /rpt_family="AluY"
 13978..14187
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 15639..15660
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 17854..18011
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 18451..18471
 /rpt_family="AT rich"
 18492..18784
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 complement(18894..19002)
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 complement(19242..19488)
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 20030..20162
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 20479..20500
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 21057..21245
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 21249..21539
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 21654..21707
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Query Match

Best Local Similarity

5.8%; Score 57.8; DB 9; Length 99268;
 67.9%; Pred. No. 0.0032;

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Matches 112; Conservative 0; Mismatches 47; Indels 6; Gaps 2;
QY 829 AGGTGAGAGCGAAGAGCAGCGCGTAGAATGAACGTCGACGAGAGC--CAAGCAC 886
Db 65407 AGGTGAGAGCATACAAAGTAGCAGGTGTAGGATGAACAGGCTAGTACTGTATA 65466
QY 887 GGCAGAGGACTGCAGGCATCAGCGTCACCTGTCGTTATTTGGAGTTCATGCCAAATGAG 946
Db 65467 ACATGAGACTGTAGGTAATAAATTGTACT---GTATGTGGAGTTCATGCTAAATGAG 65522
QY 947 TGTGTTTTAGTCTCTTGGCCACAAAAAATAAAAAAAAAAAAAA 991
Db 65523 TAGATTTTAGTCTCTTGGCCACAAACAAATACTTAAGGAA 65567

RESULT 38
AL136374
LOCUS Human DNA sequence from clone RP1-244G5 on chromosome 1q24.3-25.3,
DEFINITION complete sequence.
ACCESSION AL136374
VERSION AL136374.4 GI:8918204
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 119853)
Kimberley,A.
Direct Submission
Submitted (17-JUL-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK E-mail enquiries: humquery@sanger.ac.uk
requests: clonerequest@sanger.ac.uk
On Jul 5, 2000 this sequence version replaced gi:8176662.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit sequences with
only a small overlap as described above.
This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated human repeat sequence elements (e.g. Alu). Where the
sequence is ambiguous, there is an annotation using the 'unsure'
feature key.
The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em:, EMBL; Sw:, SWISSPROT; Tr:, TrEMBL; Wp:, WormPEP; Information
on the WormPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome 1, constructed by the Sanger Centre Chromosome 1 Mapping
Group. Further information can be found at
http://www.sanger.ac.uk/HGP/Chr1
RP1-244G5 is from the library RP1-1 constructed at the Roswell
Park Cancer Institute by the group of Pieter de Jong. For further
details see http://bacpac.med.buffalo.edu/
VECTOR: pCYPAC2
This sequence is the entire insert of clone RP1-244G5.
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/misc_feature
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repeat_region
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/note="MLT1B repeat: matches 1..390 of consensus"
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repeat_region
5284..5483
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8076..8276
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8458..8494
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repeat_region
13837..13892
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repeat_region
14392..15214
/note="MIR45B repeat: matches 1..835 of consensus"
repeat_region
16024..16224
/note="MIR repeat: matches 34..248 of consensus"
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16428..16737
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complement(17098..17761)
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misc_feature
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complement(18332..19056)
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complement(18876..19056)
/note="match: STS: Em:? Em:HSPE58B10"
18880..19060
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20198..20725
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20726..20992
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21234..21886
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22078..22430
/note="LTR16C repeat: matches 3..385 of consensus"
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22651..23063
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23449..23783
/note="MER2 repeat: matches 1..345 of consensus"

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/mol_type="genomic DNA"
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/chromosome="1"
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ORIGIN

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Best Local Similarity 56.8%; Pred. No. 0.0032;
Matches 107; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

QY      800 GTTTGGGGGTTTCAGGATAGGGAATGGGAGGTCAGAGCAGCAGCAAGCAGCAGCCATGTA 859
Db      67409 GTTGGGGTTAGAGAGGAATGAGAAATTAGTTCAGAGGATGCAAGTAGTGCATATATG 67468

QY      860 GAATGACCGTCAGAGAGCCAGCAGCGCAGAGGACTCCAGGCCATCAGCGTGCACCTGT 919
Db      67469 GGATGACCAAGTCTAGAGATCTAATAAACAACATAGGGCTATAGTTACTAAAAATTGTA 67528

QY      920 TCGATTTCGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTCCACCAAAAAA 979
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QY      980 AAAAAAAA 988
Db      67589 CCAGAAAAA 67597

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RESULT 40
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LOCUS      Homo sapiens chromosome 1 clone RP11-2619 map 1, WORKING DRAFT
DEFINITION      SEQUENCE, 15 unordered pieces.
ACCESSION      AC025375
VERSION      AC025375.3 GI:8077072
KEYWORDS      HTG; HTGS_PHASE1; HTGS_DRAFT.
SOURCE      Homo sapiens (human)
ORGANISM      Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
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Birren,B., Linton,L., Nusbaum,C., Lander,E., Abrahams,H., Allen,N.,
Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G.,
Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
Klein,J., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J.,
Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
Meldrum,J., Meneus,P., Mihova,T., Miranda,C., Mlenga,V., Morrow,J.,
Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pieani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tefaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,
Vassiliev,H., Viel,R., Vo.A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.
2 (bases 1 to 156857)
Birren,B., Linton,L., Nusbaum,C., Lander,E., Abrahams,H., Allen,N.,
Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
Boguslavskiy,L., Boukhgalter,B., Brown,A., Burkett,G.,
Campopiano,A., Castle,A., Choepel,Y., Colangelo,M., Collins,S.,
Collamore,A., Cooke,P., DeArelano,K., Dewar,K., Diaz,J.S.,
Dodge,S., Domino,M., Doyle,M., Ferreira,P., FitzHugh,W., Gage,D.,
Galagan,J., Gardyna,S., Ginde,S., Goyette,M., Graham,L.,
Grand-Pierre,N., Grant,G., Hagos,B., Heaford,A., Horton,L.,
Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kann,L., Karatas,A.,
Klein,J., LaRocque,K., Lamazares,R., Landers,T., Lehoczy,J.,
Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
Meldrum,J., Meneus,P., Mihova,T., Miranda,C., Mlenga,V., Morrow,J.,
Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
O'Neill,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
Pieani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
Stange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
Tefaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,
Vassiliev,H., Viel,R., Vo.A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
Young,G., Zainoun,J., Zimmer,A. and Zody,M.
Direct Submission
Submitted (08-MAR-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On May 25, 2000 this sequence version replaced gi:7328815.
All repeats were identified using RepeatMasker:

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Smit, A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIBR
Web site: http://www-seq.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L7958
Center clone name: 26119
----- Summary Statistics
Sequencing vector: M13; M7815; 100% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 149821 bases at least Q40
Consensus quality: 152963 bases at least Q30
Consensus quality: 154329 bases at least Q20
Insert size: 154000; agarose-fp
Insert size: 155457; sum-of-contigs
Quality coverage: 5.0 in Q20 bases; agarose-fp
Quality coverage: 5.0 in Q20 bases; sum-of-contigs
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* NOTE: This is a 'working draft' sequence. It currently
* consists of 15 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.

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1 778: contig of 778 bp in length
779 878: gap of 100 bp
879 1537: contig of 659 bp in length
1538 1637: gap of 100 bp
1638 4250: contig of 2613 bp in length
4251 4350: gap of 100 bp
4351 7895: contig of 3545 bp in length
7896 7995: gap of 100 bp
7996 11871: contig of 3876 bp in length
11872 11971: gap of 100 bp
11972 16256: contig of 4285 bp in length
16257 16356: gap of 100 bp
16357 21822: contig of 5466 bp in length
21823 21922: gap of 100 bp
21923 28321: contig of 6399 bp in length
28322 28421: gap of 100 bp
28422 36684: contig of 8263 bp in length
36685 36785: gap of 100 bp
36786 47214: contig of 10430 bp in length
47215 47314: gap of 100 bp
47315 59012: contig of 11698 bp in length
59013 59112: gap of 100 bp
59113 70244: contig of 11132 bp in length
70245 70344: gap of 100 bp
70345 86759: contig of 16415 bp in length
86760 86859: gap of 100 bp
86860 111698: contig of 24839 bp in length
111699 111799: gap of 100 bp
111799 156857: contig of 45059 bp in length.
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47296 a 31366 c 29768 g 47025 t 1402 others
ORIGIN

Query Match          5.8%; Score 57.8; DB 2; Length 156857;
Best Local Similarity 56.6%; Pred.No. 0.0032;
Matches 107; Conservative 0; Mismatches 82; Indels 0; Gaps 0;

Qy 800 GTTTGGGGGTTTCAGGATAGGAGTGGGGAGGTCAGAGGAGCGCAAGCAGCAGCAGTGA 859
Db 52955 GTTGGGGGTAGAGAGGAATCCAGAAATAGGTCAGAGGATGCGAAGTAGCATATATGTG 53014

Qy 860 GAATGACCGTCCAGAGAGCCAGCAGCGCAGAGACTGCGAGGCCATCAGCGTCACGTGT 919
Db 53015 GGATGACCAAGCTAGAGATCTAATAACACATAGGCGCTATAGTTACTAAAAATGTGA 53074

Qy 920 TCGTATTGGAGTTTCATGCAGAAATGAGTGTCTTTAGTGTCTTCCACAAAAA 979
Db 53075 CTGTATTAGATTTATGCTTAAATAGTAGATTTCAGCTGTTCTGCCAAAAACAAA 53134

Qy 980 AAAAAAAAA 988
Db 53135 CCAGAAAAA 53143

RESULT 41
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ACCESSION AC079289
VERSION HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_FULLTOP.
KEYWORDS Homo sapiens (human)
SOURCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Cararrhini; Hominidae; Homo.
REFERENCE 1 (bases 1 to 161940)
AUTHORS Birren,B., Nusbaum,C. and Lander,E.
TITLE Homo sapiens chromosome 17, clone RP11-600J16
JOURNAL Unpublished
REFERENCE 2 (bases 1 to 161940)
AUTHORS Birren,B., Linton,L., Nusbaum,C., Lander,E., Abraham,H., Allen,N.,
Anderson,S., Barna,N., Bastien,V., Beda,F., Boguslavsky,L.,
Boukhgalter,B., Brown,A., Burkett,G., Campopiano,A., Castle,A.,

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Choepe,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,P.,
DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S., Ferreira,P.,
Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S., Ginde,S., Goyette,M.,
Graham,L., Grand-Pierre,N., Hagos,B., Heatford,A., Horton,L.,
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Lamazares,R., Landers,T., Lehoczy,J., Levine,R., Lieu,C., Liu,G.,
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McPheeters,R., Meldrim,J., Meneus,L., Mihova,T., Mlenka,V.,
Morrow,J., Murphy,T., Naylor,J., Norman,C.H., O'Connor,T.,
O'Donnell,P., O'Neil,D., Oliver,T.M., Oliver,J., Peterson,K.,
Pierre,N., Pisan,C., Pollara,V., Raymond,C., Rieback,M., Riley,R.,
Rogov,P., Rothman,D., Roy,A., Santos,R., Schauer,S., Severy,P.,
Sougez,C., Spencer,B., Stange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Tirrell,A., Travers,M., Trigglio,J., Vassiliev,H., Viel,R., Vo,A.,
Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G., Zainoun,J.,
Zimmer,A. and Zody,M.
Direct Submission
Submitted (26-AUG-2000) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
3 (bases 1 to 161940)
Birren,B., Nusbaum,C., Lander,E., Ali,A., Allen,N., Anderson,S.,
Barna,N., Bastien,V., Bloom,T., Boguslavsky,L., Boukhgalter,B.,
Camarata,J., Chang,J., Chazaro,B., Choepe,Y., Collymore,A.,
Cook,A., Cooke,P., DeArellano,K., Dewar,K., Diaz,J.S., Dodge,S.,
Faro,S., Ferreira,P., Fitzgerald,M., Gage,D., Galagan,J.,
Gardyna,S., Gord,S., Graham,L., Grand-Pierre,N., Hagos,B.,
Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C., Kamat,A.,
Karatas,A., Kells,C., Landers,T., Levine,R., Lindblad-Toh,K.,
Liu,G., MacLean,C., Macdonald,P., Major,J., Matthews,C.,
McCarthy,M., Meldrim,J., Meneus,L., Mihova,T., Mlenka,V.,
Murphy,T., Naylor,J., Nguyen,C., Nicol,R., Norbu,C., Norman,C.H.,
O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J., Peterson,K.,
Phunkhang,P., Pierre,N., Raymond,C., Retta,R., Rise,C., Rogov,P.,
Roman,J., Roy,A., Schauer,S., Schuback,R., Seaman,S., Severy,P.,
Smith,C., Spencer,B., Stange-Thomann,N., Stojanovic,N., Talamas,J.,
Tesfaye,S., Theodore,J., Topham,K., Travers,M., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Young,G., Zainoun,J.,
Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (20-AUG-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
On Aug 20, 2002 this sequence version replaced gi:9929721.
All repeats were identified using RepeatMasker:
Smit,A.F.A. & Green, P. (1996-1997)
http://ftp.genome.washington.edu/RM/RepeatMasker.html
----- Genome Center
Center: Whitehead Institute/ MIT Center for Genome Research
Center code: WIER
Web site: http://www-seg.wi.mit.edu
Contact: sequence_submissions@genome.wi.mit.edu
----- Project Information
Center project name: L10440
Center clone name: 600 J 16
----- Summary Statistics
Sequencing vector: M13; M77815; 27% of reads
Chemistry: Dye-terminator Big Dye; 100% of reads
Assembly program: Phrap; version 0.960731
Consensus quality: 159446 bases at least Q40
Consensus quality: 160437 bases at least Q30
Consensus quality: 160863 bases at least Q20
Insert size: 157000; agarose-fp
Insert size: 161340; sum-of-contigs
Quality coverage: 11.9 in Q20 bases; agarose-fp
Quality coverage: 11.6 in Q2.
* NOTE: This is a 'working draft' sequence. It currently
* consists of 7 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will

```


clone sequenced to the right is RP11-642E20, 2000 bp overlap.
Actual start of this clone is at base position 97704 of RP11-35D5.

FEATURES

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1..50630
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
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/map="4"
/clone="RP11-400H10"
151..630
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repeat_region
1007..1305
/rpt_family="Alu"
repeat_region
2780..3061
/rpt_family="L2"
repeat_region
3241..3545
/rpt_family="L2"
repeat_region
4683..6142
/rpt_family="L1"
repeat_region
6143..6446
/rpt_family="Alu"
repeat_region
6447..7296
/rpt_family="L1"
repeat_region
7297..7669
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repeat_region
7670..8151
/rpt_family="L1"
repeat_region
8411..8521
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repeat_region
8732..8767
/rpt_family="AT-rich"
repeat_region
10181..10668
/rpt_family="L2"
repeat_region
10669..10936
/rpt_family="Alu"
repeat_region
10937..11264
/rpt_family="L2"
repeat_region
12145..12226
/rpt_family="L2"
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13486..13686
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repeat_region
14333..14356
/rpt_family="(CAAAA)n"
repeat_region
14395..14548
/rpt_family="MIR"
repeat_region
14678..14849
/rpt_family="MIR"
repeat_region
15282..15369
/rpt_family="L2"
repeat_region
15487..15510
/rpt_family="AT-rich"
repeat_region
15511..15572
/rpt_family="Alu"
repeat_region
15607..15753
/rpt_family="L2"
repeat_region
15982..16156
/rpt_family="L1"
repeat_region
16164..16390
/rpt_family="MaLR"
repeat_region
16406..16465
/rpt_family="MER2_type"
repeat_region
16466..18051
/rpt_family="L1"
repeat_region
18052..18369
/rpt_family="Alu"
repeat_region
18370..18939
/rpt_family="L1"
repeat_region
18960..19089
/rpt_family="L1"
repeat_region
19231..19323
/rpt_family="L1"

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repeat_region 19322..20169
/rpt_family="L1"
repeat_region 20170..20479
/rpt_family="Alu"
repeat_region 20480..20630
/rpt_family="L1"
repeat_region 20800..21207
/rpt_family="MaLR"
repeat_region 21687..21731
/rpt_family="AT-rich"
repeat_region 21782..21840
/rpt_family="L2"
repeat_region 23361..23671
/rpt_family="Alu"
repeat_region 24503..24525
/rpt_family="AT-rich"
repeat_region 26923..27223
/rpt_family="Alu"
repeat_region 27248..27504
/rpt_family="Alu"
repeat_region 27930..27951
/rpt_family="(T)n"
repeat_region 28083..28152
/rpt_family="Mariner"
repeat_region 28172..28215
/rpt_family="AT-rich"
repeat_region 29024..29147
/rpt_family="L2"
repeat_region 29148..29264
/rpt_family="(TA)n"
repeat_region 29620..29682
/rpt_family="L2"
repeat_region 30174..30291
/rpt_family="L2"
repeat_region 30570..30623
/rpt_family="CRI"
repeat_region 30632..30692
/rpt_family="MER1_type"
repeat_region 30776..30821
/rpt_family="AT-rich"
repeat_region 31291..31420
/rpt_family="L1"
repeat_region 33262..33396
/rpt_family="MIR"

Query Match      5.8%; Score 57.4; DB 9; Length 50630;
Best Local Similarity 63.1%; Pred.No. 0.0041;
Matches 123; Conservative 0; Mismatches 66; Indels 6; Gaps 2;

QY 793 AAGAGGAGTTGGGGGTTTCAGGATAGGGAATGGGAGGTTCAGAGGACGCAAGCAGCAG 852
Db 4996 AATGGTGGTTATAGGGGCAATTAATGGGAGATGTTCAGCCAGAGGATACAAAGTAGCAG 4937

QY 853 CCATGTAGATGAACCGTCCAGAGAGCCAA--GCACGGCAGAGGACTGCGGCCATCAGC 910
Db 4936 ATATGTAGGATGGAAAGTCTGGAGATCTAATGTACAAACATGAGGACTATAGGTAATAAAA 4877

QY 911 GTGCACTGTTGCTATTGTTGGAGTTTCATGCAAAATGAGTGTTTGTAGCTGCTTGGCACA 970
Db 4876 TTGTA-----TTCTATGTAGGATTCATGCTAAATGATGATAGTATTGCTGTGGCACA 4821

QY 971 AAAAAAAAAAAAAA 985
Db 4820 AAAAAACAACAAAAA 4806

RESULT 43
AL357129
LOCUS
DEFINITION Human DNA sequence from clone RP11-114A21 on chromosome
Xq21.31-22.1 Contains ESTs, STSs and GSSs, complete sequence.
ACCESSION AL357129
VERSION AL357129.11 GI:10186549

```


KEYWORDS

SOURCE Homo sapiens (human)

ORGANISM

Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 145380)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

HTG.
Submitted (19-DEC-2000) Sanger Centre, Hinxton, Cambridgeshire,
CB10 1SA, UK. E-mail enquiries: humquery@sanger.ac.uk
requests: clonerquest@sanger.ac.uk
On Sep 19, 2000 this sequence version replaced gi:9909073.
During sequence assembly data is compared from overlapping clones.
Where differences are found these are annotated as variations
together with a note of the overlapping clone name. Note that the
variation annotation may not be found in the sequence submission
corresponding to the overlapping clone, as we submit submissi-
only a small overlap as described above.

The following abbreviations are used to associate primary accession
numbers given in the feature table with their source databases:
Em:, EMBL; Sw:, SWISSPROT; Tr:, TREMBL; Mp:, WORMPEP; Information
on the WORMPEP database can be found at
http://www.sanger.ac.uk/Projects/C_elegans/wormpep This sequence
was generated from part of bacterial clone contigs of human
chromosome X, constructed by the Sanger Centre Chromosome X Mapping
Group. Further information can be found at
<http://www.sanger.ac.uk/HGP/ChrX>

This sequence has been finished according to sequence map criteria
as follows. An attempt is made to resolve all sequencing problems,
such as compressions and repeats, but not necessarily within known
annotated repeat sequence elements. Where the sequence is
ambiguous, there is an annotation using the 'unsure' feature key.
RP11-114A21 is from the library RPCI-11.1 constructed by the group
of Pieter de Jong. For further details see
<http://www.chori.org/bacpac/home.htm>

FEATURES

source

Location/Qualifiers

1..145380
/organism="Homo sapiens"
/mol_type="Genomic DNA"
/db_xref="taxon:9606"
/chromosome="X"
/map="q21.31-22.1"
/clone="RP11-114A21"
/clone_lib="RPCI-11.1"

repeat_region

1..71
/note="L1P3 repeat: matches 10..80 of consensus"

misc_feature

20..685
/note="match: GSS: Em:AQ343933"

repeat_region

31..79
/note="L1P repeat: matches 4..52 of consensus"

repeat_region

450..492
/note="MER63 repeat: matches 2..44 of consensus"

repeat_region

847..993
/note="MER20 repeat: matches 29..174 of consensus"

repeat_region

1591..1614
/note="12 copies 2 mer to 95% conserved"

repeat_region

3441..3588
/note="2 copies 74 mer 92% conserved"

repeat_region

3742..3801
/note="L2 repeat: matches 2643..2701 of consensus"

repeat_region

4195..4440
/note="MLT1A1 repeat: matches 118..365 of consensus"

repeat_region

4442..4495
/note="HERV17 repeat: matches 1..54 of consensus"

repeat_region

4496..5265
/note="LTR17 repeat: matches 1..780 of consensus"

repeat_region

5271..6471
/note="L1PBA repeat: matches 1037..2339 of consensus"

repeat_region

6464..6757
/note="L1PBA repeat: matches -712..-411 of consensus"

repeat_region

6786..7118
/note="L1PBA repeat: matches -176..-1104 of consensus"

repeat_region

7770..8014
/note="MLT1H repeat: matches 38..289 of consensus"

repeat_region

8037..8234
/note="MLT1H repeat: matches 410..547 of consensus"

repeat_region

8310..8468
/note="L2 repeat: matches 2563..2723 of consensus"

repeat_region

8484..8639
/note="L2 repeat: matches 1060..1224 of consensus"

repeat_region

9501..10717
/note="L1P3 repeat: matches 4..924 of consensus"

repeat_region

10749..11059
/note="AluY repeat: matches 1..311 of consensus"

repeat_region

11101..13160
/note="L1P repeat: matches 948..2998 of consensus"

repeat_region

13159..15022
/note="L1PBA repeat: matches 4290..6162 of consensus"

repeat_region

15671..15793
/note="L2 repeat: matches 2577..2710 of consensus"

repeat_region

16891..17018
/note="MIR repeat: matches 14..144 of consensus"

repeat_region

18274..18889
/note="L2 repeat: matches 1656..2319 of consensus"

repeat_region

18868..18953
/note="MIR repeat: matches 169..254 of consensus"

repeat_region

19189..19961
/note="L1MA2 repeat: matches 5536..6308 of consensus"

repeat_region

19962..21580
/note="L1P13 repeat: matches 4433..6073 of consensus"

repeat_region

21585..24399
/note="L1P13 repeat: matches 1532..4323 of consensus"

repeat_region

24420..25728
/note="L1M1 repeat: matches -301..1172 of consensus"

repeat_region

25721..25834
/note="L1M1 repeat: matches -1389..-1269 of consensus"

repeat_region

26112..26285
/note="MER58A repeat: matches 45..224 of consensus"

repeat_region

26387..26742
/note="MLT1A1 repeat: matches 1..365 of consensus"

repeat_region

26857..27201
/note="L1ME repeat: matches 5514..5867 of consensus"

repeat_region

27851..27978
/note="MER47 repeat: matches 2215..2323 of consensus"

repeat_region

27979..28352
/note="THE1C repeat: matches 5..371 of consensus"

repeat_region

28353..28503
/note="MER47 repeat: matches 2058..2215 of consensus"

repeat_region

28529..28645
/note="MIR repeat: matches 119..245 of consensus"

repeat_region

28672..29528
/note="MER57-internal repeat: matches 1731..2937 of consensus"

repeat_region

29688..30062
/note="HUESR-P3b repeat: matches 2227..2560 of consensus"

repeat_region

30076..30409
/note="MER57-internal repeat: matches 1306..1630 of consensus"

misc_feature

30650..30978
/note="match: GSS: Em:B38708"

misc_feature

30654..30871
/note="match: GSS: Em:AQ589046"

repeat_region

31284..31806
/note="L1R8 repeat: matches 159..691 of consensus"

repeat_region

31816..32111
/note="ALUSx repeat: matches 4..299 of consensus"

repeat_region

32145..32281

(http://bacpac.med.buffalo.edu)

VECTOR: PRACE3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is AC021789; the clone sequenced to the right is AC021688. Actual start of this clone is at base position 1 of RP11-557N1; actual end is at base position 169202 of RP11-557N1.

The sequence H NH0557N01 contains a dinucleotide (GA) repeat from base position 29551 to 29868 where the sequence fidelity cannot be guaranteed. Assembly of the database is consistent with digest information.

FEATURES

Location/Qualifiers

source

1. .169202
/organism="Homo sapiens"
/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="2"
/map="2"

/clone="RP11-557N1"

/clone.lib="RPC1-11"

196. .300

/rpt_family="Mariner"

407. .730

/rpt_family="MER121"

891. .934

/rpt_family="AT_rich"

1399. .1428

/rpt_family="AT_rich"

1550. .1577

/rpt_family="AT_rich"

1739. .1895

/rpt_family="MIR"

3511. .3635

/rpt_family="L2"

3700. .4079

/rpt_family="ERV1"

4106. .4201

/rpt_family="L2"

4947. .5133

/rpt_family="MIR"

5960. .5996

/rpt_family="polypyrimidine"

6720. .6772

/rpt_family="MIR"

6809. .6889

/rpt_family="MIR"

7122. .7348

/rpt_family="L1"

7349. .7392

/rpt_family="GTTTG)n"

7393. .7802

/rpt_family="L1"

7803. .7856

/rpt_family="(TA)n"

7867. .7984

/rpt_family="L1"

8738. .8860

/rpt_family="MIR"

8863. .9145

/rpt_family="Alu"

9426. .9714

/rpt_family="Alu"

9715. .9758

/rpt_family="AT_rich"

9903. .9977

/rpt_family="L2"

10028. .10056

/rpt_family="(TG)n"

10404. .10689

/rpt_family="Alu"

12107. .12278

/rpt_family="MaLR"

repeat_region 13854. .13946
/rpt_family="ACHobo"
repeat_region 14020. .14183
/rpt_family="MIR"
repeat_region 14531. .14841
/rpt_family="Alu"
repeat_region 14860. .15243
/rpt_family="MER2_type"
repeat_region 16032. .16338
/rpt_family="Alu"
repeat_region 17440. .17564
/rpt_family="CR1"
repeat_region 19909. .19941
/rpt_family="AT_rich"
repeat_region 20077. .20139
/rpt_family="MIR"
repeat_region 20386. .20982
/rpt_family="ERV1"
repeat_region 21156. .21524
/rpt_family="MaLR"
repeat_region 22038. .22097
/rpt_family="MIR"
repeat_region 22230. .22331
/rpt_family="MIR"
repeat_region 24135. .24157
/rpt_family="AT_rich"
repeat_region 25901. .25968
/rpt_family="MIR"
repeat_region 27874. .28003
/rpt_family="MIR"
repeat_region 28292. .28426
/rpt_family="MIR"
repeat_region 28611. .28718
/rpt_family="L2"
repeat_region 28774. .28835
/rpt_family="(TTAAA)n"
repeat_region 29147. .29261
/rpt_family="GA-rich"
repeat_region 29285. .29567
/rpt_family="Alu"
repeat_region 29568. .29642
/rpt_family="polypurine"
repeat_region 29644. .29831
/rpt_family="GA-rich"
repeat_region 29832. .29866
/rpt_family="(GGGA)n"
repeat_region 29991. .30283
/rpt_family="Alu"
repeat_region 30686. .31803
/rpt_family="L1"
repeat_region 31804. .32105
/rpt_family="Alu"
repeat_region 32106. .32750
/rpt_family="L1"

Query Match

5.8%; Score 57.4; DB 9; Length 169202;

Best Local Similarity 64.2%; Pred No. 0.004; 61; Indels 6; Gaps 2;

Matches 120; Conservative 0; Mismatches 61; Indels 6; Gaps 2;

Qy 805 GGGGTTTCAGGATAGGGAATGGGAGGTCAGAGCAGCAAGAGCAGCCATGTAGATG 864
Db 79581 GTGGAGAGGAACTGGCGGGTAGGTACGGGATAGAAATAGCAATATGAGGATG 79640
Qy 865 AACGTCAGAGAGCC--AAGCAGGAGAGGATGCGGCCATCAGCGTGCATGTTCG 922
Db 79641 AACAGGCTGGAGACCTACAGCCCAACATACCACCTACAAGTAATAAATCGAGCT---G 79696
Qy 923 TATTGGAGTTTCATGCAAAATGAGTGTGTTTACGCTCTTGCACAAAAAATAA 982
Db 79697 TATTGGAGTTTCATGCTGAATGGCTAGATTTAGCTCTTGTCCACAAAAAATAA 79756
Qy 983 AAAAAA 989
|||

```

Db      79757 TGAATAA 79763

RESULT 45
AC145171/c
LOCUS
DEFINITION Homo sapiens chromosome 16 clone RP11-349F16, WORKING DRAFT
SEQUENCE, 3 unordered pieces.
ACCESSION AC145171
VERSION AC145171.1 GI:31621312
KEYWORDS HTG; HTGS_PHASE1; HTGS_DRAFT; HTGS_ACTIVEFIN.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE
AUTHORS Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
TITLE Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
JOURNAL DOE Joint Genome Institute.
REFERENCE Sequencing of Human Chromosome 16
AUTHORS Unpublished
TITLE 2 (bases 1 to 175947)
JOURNAL DOE Joint Genome Institute.
AUTHORS Direct Submission
TITLE Submitted (12-JUN-2003) Production Sequencing Facility, DOE Joint
JOURNAL Genome Institute, 2800 Mitchell Drive, Walnut Creek, CA 94598, USA
COMMENT -----Genome Center
Center: Joint Genome Institute
Center Code: JGI
Web site: http://www.jgi.doe.gov
-----
Project Information
Center Project Name: 540517
Center clone name: RPCI-11_349F16
-----
Summary Statistics
Consensus quality: 173647 bases at least Q40
Consensus bases: 174869 bases at least Q30
Consensus quality: 175338 bases at least Q20
Estimated insert size: 175000; agarose-fp estimation
Quality coverage: 9.21 in Q20 bases; agarose-fp estimation
Quality coverage: 9.17 in Q20 bases; sum-of-contigs estimation.
NOTE: This is a 'working draft' sequence. It currently
* consists of 3 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
*
* 1 45369; contig of 45369 bp in length
* 45370 45469; gap of unknown length
* 45470 94891; contig of 49422 bp in length
* 94892 175947; gap of unknown length
* 94992 175947; contig of 80956 bp in length.
* Location/Qualifiers
* 1..175947
* /organism="Homo sapiens"
* /mol_type="genomic DNA"
* /db_xref="taxon:9606"
* /chromosome="16"
* /clone="RP11-349F16"
* /clone_lib="RPCI human BAC library 11"
* 54782 a 31550 c 31265 g 58150 t 200 others

BASE COUNT 54782 a 31550 c 31265 g 58150 t 200 others
ORIGIN

Query Match 5.8%; Score 57.4; DB 2; Length 175947;
Best Local Similarity 55.9%; Pred. No. 0.004;
Matches 109; Conservative 0; Mismatches 86; Indels 0; Gaps 0;

QY 795 GAGAGATTGGGGTTTCAGATAGGAATGGGAGTCTCAGAGACCAAGCAGCAGCC 854
Db 39798 GTGGGGATGGAGTGGGACTAAATGAGGAGAGGTAGGTTCAGATTGCAAGTAGCAGAT 39739

```

```

QY 855 ATGTAGATGAACCGTCCAGAGAGCCAGCAGCGAGTCTGAGCCATCAGCGTGC 914
Db 39738 AAGTAGGATGAACAGCTCAGAGATCTAATGTACAATATGAGAACTATAGCTTTAAAAA 39679
QY 915 ACTGTTGCTATTGAGTTCATGCAAAATGAGTGTGTTTGTAGCTGCTCTTGCCACAAAAA 974
Db 39678 ATTGTATTGATTGGGTTTCCTGTTAAATGAGTACGTTTGTAGCTGCTCTTGCTACAAAAT 39619
QY 975 AAAAAAARAAAAA 989
Db 39618 CAACACAAAGAAAA 39604

RESULT 46
AC013408
LOCUS
DEFINITION Homo sapiens BAC clone RP11-469G18 from 2, complete sequence.
ACCESSION AC013408
VERSION AC013408.7 GI:15920142
KEYWORDS HTG.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
TITLE 1 (bases 1 to 151841)
JOURNAL Sulston, J.E. and Waterston, R.
REFERENCE Toward a complete human genome sequence
AUTHORS Genome Res. 8 (11), 1097-1108 (1998)
TITLE 99063792
JOURNAL MEDLINE
REFERENCE 9847074
AUTHORS 2 (bases 1 to 151841)
TITLE Holmes, A., Hawkins, M. and Kozlowski, A.
JOURNAL The sequence of Homo sapiens BAC clone RP11-469G18
REFERENCE Unpublished (2001)
AUTHORS 3 (bases 1 to 151841)
TITLE Waterston, R.H.
JOURNAL Direct Submission
REFERENCE Submitted (09-NOV-1999) Genome Sequencing Center, Washington
AUTHORS University School of Medicine, 4444 Forest Park Parkway, St. Louis,
TITLE MO 63108, USA
JOURNAL 4 (bases 1 to 151841)
REFERENCE Waterston, R.H.
AUTHORS Direct Submission
TITLE Submitted (04-OCT-2001) Genome Sequencing Center, Washington
JOURNAL University School of Medicine, 4444 Forest Park Parkway, St. Louis,
REFERENCE MO 63108, USA
AUTHORS 5 (bases 1 to 151841)
TITLE Waterston, R.
JOURNAL Direct Submission
REFERENCE Submitted (09-JAN-2002) Department of Genetics, Washington
AUTHORS University, 4444 Forest Park Avenue, St. Louis, Missouri 63108, USA
TITLE On Oct 4, 2001 this sequence version replaced gi:13992768.
JOURNAL COMMENT
Center: Washington University Genome Sequencing Center
Center code: WUGSC
Web site: http://genome.wustl.edu/gsc
Contact: sapiens@watson.wustl.edu
Summary Statistics
Center project name: H_NF0469G18

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NOTICE: This sequence may not represent the entire insert of this clone. It may be shorter because we only sequence overlapping clone sections once, or longer because we provide a small overlap between neighboring data submissions.

This sequence was finished as follows unless otherwise noted: all regions were double stranded, sequenced with an alternate chemistry, or covered by high quality data (i.e., phred quality >= 30); an attempt was made to resolve all sequencing problems, such as compressions and repeats; all regions were covered by sequence from more than one subclone; and the assembly was confirmed by restriction digest.

MAPPING INFORMATION:

Mapping information for this clone was provided by Dr. John D. McPherson, Department of Genetics, Washington University, St. Louis MO. For additional information about the map position of this sequence, see <http://genome.wustl.edu/gsc>

SOURCE INFORMATION:

The RPCI-11 human BAC library was made from the blood of one male donor, as described by Osogawa, K., Woon, P.Y., Zhao, B., Frengen, E., Tateno, M., Catanese, J.J., and de Jong, P.J. (1998) An improved approach for construction of bacterial artificial chromosome libraries. *Genomics* 51:1-8. The clone may be obtained either from Research Genetics, Inc. (<http://www.resgen.com>) or Pieter de Jong and coworkers at the Roswell Park Cancer Institute (<http://bacpac.med.buffalo.edu>)

VECTOR: pBACe3.6

NEIGHBORING SEQUENCE INFORMATION:

The clone sequenced to the left is RP11-90C9, 2000 bp overlap; the clone sequenced to the right is RP11-22L19, 2000 bp overlap. Actual start of this clone is at base position 1 of RP11-469G18; actual end is at base position 14496 of RP11-22L19.

Data from AC079146 was used to finish this clone, AC013408.

Polymorphisms have been identified between AC079146 and AC013408.

FEATURES

source

Location/Qualifiers

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Query Match

Best Local Similarity 56.0%; Score 56.6; DB 9; Length 151841;

Matches 107; Conservative 0; Mismatches 84; Indels 0; Gaps 0;

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Db      124652  ATAGCAGGTATGTAGAAATGAACAGTCTGAACATCTAAATAAATGATTAACCATAGTTAA 124711
QY      906  TCAGCTGCACCTGTTCTGTTATTTGGAGTTCATGCAAAATGAGTGTGTTTATGCTGCTCTTG 965
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RESULT 47
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LOCUS      173270 bp      DNA      linear      PRI 05-FEB-2003
DEFINITION Homo sapiens chromosome 8, clone RP11-599P7, complete sequence.
ACCESSION AC025522
VERSION   AC025522.7  GI:28209714
KEYWORDS  HTG.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
  1 (bases 1 to 173270)
  Birren,B., Nussbaum,C. and Lander E.
  Homo sapiens chromosome 8, clone RP11-599P7
  Unpublished
  2 (bases 1 to 173270)
  Birren,B., Linton,L., Nussbaum,C., Lander,E., Abraham,H., Allen,N.,
  Anderson,S., Baldwin,J., Barna,N., Bastien,V., Beda,F.,
  Boguslavsky,L., Boukhgalter,B., Brown,A., Burkett,G.,
  Campopiano,A., Castile,A., Choepel,Y., Colangelo,M., Collins,S.,
  Collymore,A., Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S.,
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  Howland,J.C., Iliev,I., Johnson,R., Jones,C., Kaur,L., Karatas,A.,
  Klein,J., LaRoque,K., Lamazares,R., Landers,T., Lehoczy,J.,
  Levine,R., Lieu,C., Liu,G., Locke,K., Macdonald,P., Marquis,N.,
  McCarthy,M., McEwan,P., McGurk,A., McKernan,K., McPheeters,R.,
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  Murphy,T., Naylor,J., Norman,C.H., O'Connor,T., O'Donnell,P.,
  O'Neil,D., Oliver,T.M., Oliver,J., Peterson,K., Pierre,N.,
  Pisani,C., Pollara,V., Raymond,C., Riley,R., Rogov,P., Rothman,D.,
  Roy,A., Santos,R., Schauer,S., Severy,P., Spencer,B.,
  Strange-Thomann,N., Stojanovic,N., Subramanian,A., Talamas,J.,
  Tesfaye,S., Theodore,J., Tirrell,A., Travers,M., Trigilio,J.,
  Vassiliev,H., Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J.,
  Young,G., Zainoun,J., Zimmer,A. and Zody,M.
  Direct Submission
  Submitted (09-MAR-2000) Whitehead Institute/MIT Center for Genome
  Research, 320 Charles Street, Cambridge, MA 02141, USA
  3 (bases 1 to 173270)
  Birren,B., Linton,L., Nussbaum,C., Lander,E., Ali,A., Allen,N.,
  Anderson,S., Barna,N., Bastien,V., Boguslavsky,L., Boukhgalter,B.,
  Brown,A., Camarata,J., Campopiano,A., Chang,J., Chazaro,B.,
  Choepel,Y., Colangelo,M., Collins,S., Collymore,A., Cooke,A.,
  Cooke,P., Dearellano,K., Dewar,K., Diaz,J.S., Dodge,S., Faro,S.,
  Ferreira,P., Fitzhugh,W., Gage,D., Galagan,J., Gardyna,S.,
  Ginde,S., Gord,S., Goyette,M., Graham,L., Grand-Pierre,N.,
  Hagos,B., Horton,L., Hulme,W., Iliev,I., Johnson,R., Jones,C.,
  Kamat,A., Karatas,A., Kells,C., LaRoque,K., Lamazares,R.,
  Landers,T., Lehoczy,J., Levine,R., Liu,G., Maclean,C.,
  Macdonald,P., Major,J., Marquis,N., Matthews,C., McCarthy,M.,
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  Norman,C.H., O'Connor,T., O'Donnell,P., O'Neil,D., Oliver,J.,

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Peterson,K., Phunkhang,P., Pierre,N., Pollara,V., Raymond,C.,
Retta,R., Rieback,M., Riley,R., Rise,C., Rogov,P., Roman,J.,
Rosetti,M., Roy,A., Santos,R., Schauer,S., Schuback,R., Seaman,S.,
Severy,P., Spencer,B., Strange-Thomann,N., Stojanovic,N.,
Strauss,N., Subramanian,A., Talamas,J., Tesfaye,S., Theodore,J.,
Topham,K., Travers,M., Travis,N., Trigilio,J., Vassiliev,H.,
Viel,R., Vo,A., Wilson,B., Wu,X., Wyman,D., Ye,W.J., Young,G.,
Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
Direct Submission
Submitted (12-FEB-2002) Whitehead Institute/MIT Center for Genome
Research, 320 Charles Street, Cambridge, MA 02141, USA
4 (bases 1 to 173270)
  Birren,B., Nussbaum,C., Lander,E., Abouelleil,A., Allen,N.,
  Anderson,S., Arachchi,H.M., Barna,N., Bastien,V., Bloom,F.,
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  Wyman,D., Young,G., Zainoun,J., Zembek,L., Zimmer,A. and Zody,M.
  Direct Submission
  Submitted (05-FEB-2003) Whitehead Institute/MIT Center for Genome
  Research, 320 Charles Street, Cambridge, MA 02141, USA
  On Feb 5, 2003 this sequence version replaced gi:18652546.
  All repeats were identified using RepeatMasker:
  Smit, A.F.A. & Green, P. (1996-1997)
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  ----- Genome Center
  Center: Whitehead Institute/ MIT Center for Genome Research
  Web site: http://www-seq.wi.mit.edu
  Contact: sequence_submissions@genome.wi.mit.edu
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DEFINITION		AC016256	
ACCESSION		AC016256.19	GI:14670071
VERSION		HTG.	
KEYWORDS		Homo sapiens (human)	
SOURCE		Homo sapiens	
ORGANISM		Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	
REFERENCE		Mammalia; Euthera; Primates; Catarrhini; Homnidae; Homo.	
AUTHORS		Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-Osman,F.R., Allen,C.,	
		Alsbrooks,S.L., Amaratunge,H.C., Are,J.-R., Banks,T., Barbarella,J.,	
		Benton,J., Binage,K., Blankenburg,K., Bonnin,D., Bouck,J.J.,	
		Bowie,S., Brieva,M., Brown,E., Brown,M., Bryant,N.P., Buhay,C.,	
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		Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,	
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		Zorrilla,S., Zucherlapati,R. and Gibbs.R.	

TITLE

Direct Submission

Unpublished

2 (bases 1 to 164405)

REFERENCE

Worley, K.C.

Direct Submission

Submitted (24-NOV-1999) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

3 (bases 1 to 164405)

REFERENCE

Worley, K.C.

Direct Submission

Submitted (11-JUL-2001) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

4 (bases 1 to 164405)

REFERENCE

Worley, K.C.

Direct Submission

Submitted (25-JUL-2001) Human Genome Sequencing Center, Department

of Molecular and Human Genetics, Baylor College of Medicine, One

Baylor Plaza, Houston, TX 77030, USA

COMMENT

On Jul 11, 2001 this sequence version replaced gi:12656764.

INFORMATION: <http://www.hgsc.bcm.tmc.edu/> or email

gc-help@bcm.tmc.edu

CLONE LENGTH: This sequence does not necessarily represent the entire insert of this clone. Overlapping regions of clones are only sequenced and submitted once, so the sequence for the remainder of the insert may be found in the record for the adjacent clones. Overlapping clones are noted at the beginning and end of the Features listing.

ANNOTATION OF FEATURES:

STSS are identified using ePCR (Genome Res. 7:541-550) searches of a local database that includes entries from dbSTS, GDB, and local mapping efforts.

Repeats are identified using RepeatMasker (A. Smit and P. Green, unpublished.) for Human and Mouse sequences.

Genes and Region of sequence similarity are identified by BLAST (Nuc. Acids Res. 25:3389-3402) similarity (expect < 1e-34) to the EST and cDNA sequences. Genes demonstrate at least two exons flanked by consensus splice sites that maintained sequence continuity across the splice junctions. Sequences that are not identical matches are annotated as similar.

SEQUENCING READ COVERAGE: Sequencing is completed to a minimum standard of double strand coverage with a minimum of 2 clones and 2 reads with no ambiguities or 2 chemistries with a minimum of 2 clones and 3 reads with no ambiguities. If the sequence quality for a region does not meet this standard, it will be indicated in the annotation as Low Coverage.

QUALITY OF INDIVIDUAL BASES: This sequence meets stringent quality standards - estimated error rate less than 1 per 10,000 bases. Reports of lowest quality individual bases and measures of base quality are listed below. Description of the metrics can be found at URL:

<http://gc.bcm.tmc.edu:8086/quality.info/genbank.annotation.html>.

QUALSTAT-REPORT.

FEATURES

source

Location/Qualifiers

1..164405

/organism="Homo sapiens"

/mol_type="genomic DNA"

/db_xref="taxon:9606"

/chromosome="12"

/clone="RP11-98E6"

complement(1..371)

/rpt_family="MLTIF1"

3..5166

/notes="Overlaps bases 186760..191923 of clone AC087886"

/function="Overlaps with adjacent clone AC087886"

complement(439..637)

/rpt_family="L1MA8"

repeat_region

misc_feature

repeat_region

repeat_region

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2142..2281
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7015..7068
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7069..7153
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7163..7191
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/note="CA or TG"


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Matches 137; Conservative 0; Mismatches 96; Indels 3; Gaps 2;

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Db 143164 CCTTGAATATATAGATAGATAGAGAAGAAGACAGTGTGATACAGGTGGAGGGTGGTCCAAAA 143223

QY 817 TAGGGATGGGGAGGTGAGAGGACGCGAAGAGCAGAGCATGTAGAAATGAACCGTCCAGAG 876
Db 143224 ATGGGAAGATGTAGTTAAGAGGATACAAATTA--AGTTATGTAGATGAACAGTCTAGA 143281

QY 877 AGCCAACGCGGACGAGACTGCGAGGCGCATCAGCGTGCATCTGTTGATTTGAGTTTCAT 936
Db 143282 GGTCTTAGATACATATGAGGACTATATAATTAATAAATTGACT-GTGTCTGTGATTCAT 143340

QY 937 GCAAAATGAGTGTGTTTGTAGTCTCTTGTCCACAAAAAATAAAAAAATAAAAAA 992
Db 143341 GCTAATGATGTCGATTTTGTAGTCTCTTGTCCACAAAAAATAAAAAAATAAAAAA 143396

RESULT 49
AC025258/c
LOCUS
DEFINITION
Homo sapiens chromosome 12 clone RP11-398017, WORKING DRAFT
ACCESSION
AC025258.15 GI:14547352
VERSION
HTG; HTGS PHASE1; HTGS_DRAFT.
KEYWORDS
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
Muzny,D.M., Adams,C., Adio-Oduola,B., Ali-oaman,F.R., Allen,C.,
Alsbrooks,S.L., Amarutunge,H.C., Are,J.R., Banks,T., Barbara,J.,
Benton,J., Binage,K., Blankenburg,K., Bonnin,D., Bouck,J.,
Bowles,S., Brieva,M., Brown,M., Bryant,N.P., Buhay,C.,
Burch,P., Burkett,C., Burrell,K.L., Byrd,N.C., Carron,T.F.,
Carter,M., Cavazos,S.R., Chacko,J., Chavez,D., Chen,G., Chen,R.,
Chen,Z., Chowdhry,I., Christopoulos,C., Cleveland,C.D., Cox,C.,
Coyle,M.D., Dathorne,S.R., David,R., Davila,M.L., Davis,C.,
Davy-Carroll,L., Dederich,D.A., Delaney,K.R., Delgado,O.,
Denn,A.L., Ding,Y., Dinh,H.H., Douthwaite,K.J., Draper,H.,
Dugan-Rocha,S., Durbin,K.J., Earnhart,C., Edgar,D., Edwards,C.C.,
Elhaj,C., Escotto,M., Falls,T., Ferraguto,D., Flagg,N., Ford,J.,
Foster,P., Frantz,P., Gabisi,A., Gao,J., Garcia,A., Garner,T.,
Garza,N., Gill,R., Gorrell,J.H., Guevara,W., Gunaratne,P., Hale,S.,
Hamilton,K., Harris,C., Harris,K., Hart,M., Havlak,P., Hawes,A.,
Hernandez,J., Hernandez,O., Hodgson,A., Hogue,M., Holloway,C.,
Hollins,B., Homs,F., Howard,S., Huber,J., Hulyk,S., Hume,J.,
Jackson,L.E., Jacobson,B., Jia,Y., Johnson,R., Jolivet,S.,
Joudah,S., Karlsson,E., Kureshi,A., Landry,N., Leal,B., Lewis,L.C.,
Kovach,S., Kratovic,J., Kelly,S., Khan,U., King,L., Korvah,J.,
Lewis,L., Li,J., Li,Z., Lichtarge,O., Lien,C., Liu,J., Liu,W.,
Loulsegh,H., Lozano,R.J., Lu,X., Lucier,A., Lucier,R., Luna,R.,
Ma,J., Maheshwari,M., Mapua,P., Martin,R., Martinale,A.,
Martinez,E., Massey,E., Mawhinney,E., McLeod,M.P., Meador,M.,
Mei,G., Metzker,M., Miner,G., Miner,Z., Mitchell,T., Mohabbat,K.,
Morgan,A., Morris,M., Moser,M., Neal,D., Newton,J., Newton,N.,
Nguyen,A., Nguyen,N., Nguyen,N., Nickerson,E., Nwokwenkwo,S.,
Ogih,M., Okwuonu,G., Oragunye,N., Oviedo,R., Pace,A., Payton,B.,
Peery,L., Perez,L., Peters,L., Pickens,E., Primus,E., Pu,L.I.,
Quiles,M., Ren,Y., Rives,M., Rojas,A., Rojibokan,I., Rolfe,M.,
Ruiz,S., Savery,G., Scherer,S., Scott,G., Shen,H., Shoohtari,N.,
Sisson,I., Sodergren,E., Sonaike,T., Sparks,A., Stanley,H.,
Stone,H., Sutton,A., Svatek,A., Tabor,P., Tamerisa,A., Tamerisa,K.,
Tang,H., Tansey,J., Taylor,C., Taylor,T., Telford,B., Thomas,N.,
Thomas,S., Umani,K., Vasquez,L., Vera,V., Villalon,D., Vinson,R.,
Wall,R., Wang,S., Ward-Moore,S., Warren,R., Washington,C.,
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Watlington,S., Williams,G., Williamson,A., Wleczyk,R., Wooden,S.,
Worley,K., Wu,C., Wu,Y., Wu,Y.F., Zhou,J., Zorrilla,S., Nelson,D.,
Weinstock,G. and Gibbs,R.
Direct Submission
Unpublished
2 (bases 1 to 166668)
Worley,K.C.
Direct Submission
Submitted (08-MAR-2000) Human Genome Sequencing Center, Department
of Molecular and Human Genetics, Baylor College of Medicine, One
Baylor Plaza, Houston, TX 77030, USA
On Jun 25, 2001 this sequence version replaced gi:12621178.
----- Genome Center
Center: Baylor College of Medicine
Center code: BCM
Web site: http://www.hgsc.bcm.tmc.edu/
Contact: hgsc-help@bcm.tmc.edu
----- Project Information
Center project name: HALL
Center clone name: RP11-398017
----- Summary Statistics
Sequencing vector: M13; L08821
Chemistry: Dye-primer Bodipy; 27% of reads
Chemistry: Dye-terminator Big Dye; 73% of reads
Assembly program: Phrap; version 0.990329
Consensus quality: 177529 bases at least Q40
Consensus quality: 182522 bases at least Q30
Consensus quality: 185426 bases at least Q20
Estimated insert size: 170888; sum-of-contigs estimation
Quality coverage: 0x in Q20 bases; agarose-tp estimation
Quality coverage: 6.3x in Q20 bases; sum-of-contigs estimation
-----
* NOTE: Estimated insert size may differ from sequence length
* (see http://www.hgsc.bcm.tmc.edu/docs/Genbank_draft_data.html).
* NOTE: This is a 'working draft' sequence. It currently
* consists of 2 contigs. The true order of the pieces
* is not known and their order in this sequence record is
* arbitrary. Gaps between the contigs are represented as
* runs of N, but the exact sizes of the gaps are unknown.
* This record will be updated with the finished sequence
* as soon as it is available and the accession number will
* be preserved.
* 1 149576: contig of 149576 bp in length
* 149577 149676: gap of unknown length
* 149677 166668: contig of 16992 bp in length.
* Location/Qualifiers
* 1.166668
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* /mol_type="genomic DNA"
* /db_xref="taxon:9606"
* /chromosome="12"
* /clone="RP11-398017"
BASE COUNT 58175 a 29138 c 28414 g 50819 t 122 others
ORIGIN
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Best Local Similarity 58.1%; Pred. No. 0.0069;
Matches 137; Conservative 0; Mismatches 96; Indels 3; Gaps 2;
QY 757 CCTTGGCCGCTCTACAGGAGCACCCCGCTCTGAGTGAAGAGGAGTCTTGGGGGTTTCAGGA 816
Db 111278 CCTTGAATATGATAGATAGAGAAGAAGACAGTGTGATACAGGTGGAGGGTGGTCCAAAA 111219

QY 817 TAGGGAATGGGGAGGTGAGAGGACGCGAAGAGCAGAGCATGTAGAAATGAACCGTCCAGAG 876
Db 111218 ATGGGAAGATGATCTTAAGAGGATACAAATTA--AGTTATGTAGATGAACAGTCTAGA 111161

QY 877 AGCCAACGCGGACGAGACTGCGAGGCGCATCAGCGTGCATCTGTTGATTTGAGTTTCAT 936
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QY 937 GCAAAATGAGTGTGTTTGTAGTCTCTTGTCCACAAAAAATAAAAAAATAAAAAA 992
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Db 111101 GCTAATGAGTCGATTAGTCTCTTGGCCACAAACCAAAAAAAAAAAAAAAAA 111046

RESULT 50
AC011004/c
LOCUS
DEFINITION Homo sapiens chromosome 8 clone RP11-110D15, WORKING DRAFT
AC011004
ACCESSION AC011004.12 GI:13273359
VERSION HTG; HTGS PHASE1; HTGS_DRAFT; HTGS_FUILLTOP.
KEYWORDS Homo sapiens (human)
SOURCE Homo sapiens
ORGANISM Homo sapiens
REFERENCE
AUTHORS 1 (bases 1 to 183778) Dela Rosa, M., Federspiel, N., Foreman, P., Glukhov, S., Hansen, N., Hyman, R., Mao, J., Marathe, R., Morehouse, A.J., Oefner, P., Palm, C.J., Ramirez, D. and Davis, R.W. Direct Submission
JOURNAL Submitted (29-SEP-1999) DNA Sequencing and Technology Center, Stanford University, 855 California Avenue, Palo Alto, CA 94304, USA

On Mar 11, 2001 this sequence version replaced gi:13270571.
----- Genome Center
Center: Stanford DNA Sequencing and Technology Development Center
Center code: SOSTDC
Web site: <http://sequence-www.stanford.edu/group/human/>
Contact: hum-info@sequence.stanford.edu
----- Project Information
Center project name: 666
Center clone name: RP11-110D15
----- Summary Statistics
Sequencing Vector: M13mp18; X02513
Chemistry: Dye-primer; 0% of reads
Assembly: Dye-terminator Big Dye; 100% of reads
Consensus quality: 182108 bases at least Q40
Consensus quality: 183015 bases at least Q30
Consensus quality: 183336 bases at least Q20
Insert size: 185712; agarose-fp
Quality coverage: 9.3x in Q20 bases; agarose-fp
Quality coverage: 9.4x in Q20 bases; sum-of-contigs.
NOTE: This is a 'working draft' sequence. It currently consists of 3 contigs. The true order of the pieces is not known and their order in this sequence record is arbitrary. Gaps between the contigs are represented as runs of N, but the exact sizes of the gaps are unknown. This record will be updated with the finished sequence as soon as it is available and the accession number will be preserved.

1 26915: contig of 26915 bp in length
* 26916 27015: gap of unknown length
* 27016 98873: contig of 71858 bp in length
* 98874 98973: gap of unknown length
* 98974 183778: contig of 84805 bp in length.
Location/Qualifiers
1. 183778
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/mol_type="genomic DNA"
/db_xref="taxon:9606"
/chromosome="8"
/clone="RP11-110D15"
/clone_lib="RPCI human BAC library 11"

misc_feature 1. 26915
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clone_end:SP6"
27016-.98873
misc_feature
/note="assembly_name:Contig9
clone_end:T7"
98974..183778
misc_feature
/note="assembly_name:Contig10"
201 others
BASE COUNT 50361 a 41935 c 41401 g 49880 t
ORIGIN

Query Match 5.6%; Score 56; DB 2; Length 183778;
Best Local Similarity 64.4%; Pred. No. 0.0087;
Matches 116; Conservative 0; Mismatches 60; Indels 4; Gaps 2;
QV 813 AGGATAGGGAATGGGGAGGTCTAGAGGAGCGCAAGCAGCAGCATGTAGAAATGAACCGTCC 872
DB 37638 AGGAAACGGGAGATCTAGTGTGAAGGACGACAGAGGAGATATGCAAGATGCATCGCCC 37579
QV 873 AGAGAGCCCAAGCAGCGGACAGAGACTGCGAGGCCATCAGCGTGCACCTGTTCTGATTGGAGT 932
DB 37578 AGAGACCTTAATGT-GCATGAGGACTCTCTAGTCAACCAATTGA---ATTATACCGGGAT 37523
QV 933 TCATGCAAAATGAGTGTCTTTTAGTCTCTTGGCCACAAAAAAAAAAAAAAAAAAAAAA 992
DB 37522 TTCTGCTAAGACAGTAGATTTTAGGAGCTCTTGGCATATTGAAAAAAAAAAAAAAAAAAAA 37463

Search completed: February 9, 2004, 14:16:33
Job time : 3928 secs

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 9, 2004, 13:03:52 ; Search time 2524 Seconds
(without alignments)
9552.314 Million cell updates/sec

Title: US-09-990-726-222

Perfect score: 992

Sequence: 1 ggcacgacgcaggactagg.....aaaaaaaaaaaaaaaaaaaa 992

Scoring table: IDENTITY_NUC

Gapop 10.0 , Gapext 1.0

Searched: 22781392 seqs, 12152238056 residues

Total number of hits satisfying chosen parameters: 45562784

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

EST:*

1: em_estba:*

2: em_estbum:*

3: em_estin:*

4: em_estnu:*

5: em_estov:*

6: em_estpl:*

7: em_estro:*

8: em_hic:*

9: gb_estli:*

10: gb_est2:*

11: gb_hic:*

12: gb_est3:*

13: gb_est4:*

14: gb_est5:*

15: em_estfun:*

16: em_estom:*

17: em_gss_hum:*

18: em_gss_inv:*

19: em_gss_pln:*

20: em_gss_vrt:*

21: em_gss_fun:*

22: em_gss_mam:*

23: em_gss_mus:*

24: em_gss_pro:*

25: em_gss_rod:*

26: em_gss_phg:*

27: em_gss_vrl:*

28: gb_gsel:*

29: gb_gsel2:*

Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

Result No.	Score	Query Match	Length	ID	Description
1	626.8	63.2	694	13	BX097129 BX097129
C 2	409.8	41.3	414	14	H74303 YU56C11.S1
C 3	408	41.1	437	14	N54458 YV40B02.S1
4	373.2	37.6	480	14	H73373 YU48F10.R1

5	354	35.7	418	14	H58326	H58326 Yr25c07.r1
6	349	35.2	582	14	H74302	H74302 YU56C11.R1
7	340.6	34.3	468	14	N76568	N76568 YV40B02.R1
8	320.4	32.3	996	11	AK020081	AK020081 Mus muscu
C 9	286	28.8	405	9	AI242058	AI242058 qb81d07.x
C 10	284.2	28.6	296	14	H73374	H73374 YU48F10.S1
C 11	277.2	27.9	379	14	R02548	R02548 ye8oa07.s1
C 12	274.2	27.6	450	9	AI438986	AI438986 tcb8a08.x
C 13	268.8	27.1	553	14	BY718164	BY718164 BY718164
C 14	265.2	26.7	528	4	BX529784	BX529784 RZPD Mus
C 15	261.2	26.3	384	14	H40263	H40263 YP59E11.S1
C 16	256.8	25.9	284	14	H58715	H58715 Yr25c07.s1
C 17	235.2	23.7	376	14	T80187	T80187 Yd85G03.S1
C 18	227.4	22.9	334	9	AA676653	AA676653 ZJ67E03.S
C 19	227	22.9	227	9	AI022453	AI022453 OW96H04.X
C 20	220.2	22.2	358	14	R02666	R02666 ye8oa07.r1
C 21	218.4	22.0	1161	14	W41959	W41959 mc68B10.R1
C 22	202	20.4	401	9	AA050499	AA050499 mJ20F02.F
C 23	192.2	19.4	324	14	T87491	T87491 Yd85G03.R1
C 24	182.2	18.4	467	14	W65713	W65713 me09G06.R1
C 25	175.6	17.7	218	14	N86939	N86939 L1498F Huma
C 26	174.6	17.6	364	13	BY212485	BY212485 BY212485
C 27	154	15.5	537	12	BI347002	BI347002 376394 MA
C 28	154	15.5	582	12	BI347003	BI347003 376395 MA
C 29	153.8	15.5	443	14	CB051335	CB051335 NISC_GJ23
C 30	153.8	15.5	540	14	CB051334	CB051334 NISC_GJ23
C 31	145.2	14.6	194	14	N88474	N88474 K3475F Huma
C 32	143	14.4	382	10	BB842514	BB842514 BB842514
C 33	132.6	13.4	370	10	BB843270	BB843270 BB843270
C 34	93.2	9.4	420	10	BF443482	BF443482 260980 MA
C 35	92.6	9.3	503	10	BF442092	BF442092 258654 MA
C 36	79.8	8.0	208	9	AA031034	AA031034 mJ47H05.F
C 37	57.8	5.8	483	9	AI187208	AI187208 gf28B11.X
C 38	52.8	5.3	466	9	AW794912	AW794912 RC6-UM001
C 39	52.6	5.3	493	2	HSNC077193	Bx487006 Homo sapi
C 40	51.2	5.2	606	12	BI063151	BI063151 IL3-DT011
C 41	50.8	5.1	443	13	BY560588	BY560588 BY560588
C 42	49	4.9	925	29	CNS00512P	AL053013 Drosophila
C 43	48	4.8	844	29	CNS00521P	AL056652 Drosophila
C 44	47.4	4.8	471	9	AW795001	AW795001 RC6-UM001
C 45	47.2	4.8	974	13	BX339244	BX339244 BX339244
C 46	46.4	4.7	748	29	BZ772737	BZ772737 mcv51f02
C 47	46.2	4.7	1201	13	BX356664	BX356664 BX356664
C 48	45.6	4.6	569	14	CB615918	CB615918 AMGNNUC:U
C 49	45.6	4.6	626	14	CD397038	CD397038 Gm_CK1771
C 50	45.6	4.6	935	29	CNS006XK	AL066051 Drosophila
C 51	45.4	4.6	925	29	CNS0091P	AL053013 Drosophila
C 52	45.4	4.6	1201	13	BX381961	BX381961 BX381961
C 53	45	4.5	537	14	CA442247	CA442247 UI-H-D10-
C 54	44.8	4.5	739	29	AG187401	AG187401 Pan trogl
C 55	44.6	4.5	638	14	CB242205	CB242205 UI-CF-PN0
C 56	44.2	4.5	1100	29	CNS00FSE	AL070988 Drosophila
C 57	43.4	4.4	660	13	BU928339	BU928339 AGENCOURT
C 58	43.2	4.4	420	9	AA3333438	AA3333438 S21G11 AG
C 59	43	4.3	449	9	AA186675	AA186675 ZP71C08.S
C 60	43	4.3	591	14	CB000366	CB000366 S345U G08
C 61	43	4.3	699	29	AG169292	AG169292 Pan trogl
C 62	43	4.3	940	13	BU185569	BU185569 AGENCOURT
C 63	42.6	4.3	349	28	AQ097268	AQ097268 HS 3038 B
C 64	42.6	4.3	524	28	AQ480601	AQ480601 RPCI-11-B
C 65	42.6	4.3	628	29	AG018533	AG018533 Homo sapi
C 66	42.6	4.3	638	29	AG018534	AG018534 Homo sapi
C 67	42.6	4.3	639	29	AG017811	AG017811 Homo sapi
C 68	42.6	4.3	641	29	AG017810	AG017810 Homo sapi
C 69	42.6	4.3	643	29	AG017808	AG017808 Homo sapi
C 70	42.4	4.3	699	14	CB046795	CB046795 NISC_gf06
C 71	42.2	4.3	385	9	AI478457	AI478457 tm44C09.X
C 72	42.2	4.3	469	13	BQ169652	BQ169652 WHE0961.B
C 73	42.2	4.3	1036	9	AL550172	AL550172 AL550172
C 74	42	4.2	543	9	AV597018	AV597018 AV597018
C 75	42	4.2	549	28	AQ221080	AQ221080 HS 3223 B
C 76	41.8	4.2	260	10	BF763246	BF763246 RC6-CS007
C 77	41.8	4.2	289	9	AM665077	AM665077 hi86h11.X


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QY      583  CATGCCACAGGCGCTGCCAACTTCTCTCTGCGGAGCCAGACATCGGACTGTGTCT 642
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QY      703  GTGGTCAACAGAGATGGAGGACTGCGAGGTCCTCCCTGGAGAGCCCCCATCTTGCCTTGC 762
Db      447  GTGGTCAACAGAGATGGAGGACTGCGAGGTCCTCCCTGGAGAGCCCCCATCTTGCCTTGC 506
QY      763  CGCTCTACAGGAGCACCCGCCCTCTGAGTGAAGAGAGTTTGGGGGTTTCAGATAGGCA 822
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QY      823  AT-GGGAGGTGAGGAGCGCAAGACGACGACCATGTAGATGAACCGTCCAGAGAGCA 881
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QY      882  AGCAGGCGACGAGACTGCGAGGCGCATCAGCGTGCCTGTTGCTGATTTGGAGTTTCATGCAA 941
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QY      942  ATGAGTGT 949
Db      687  ATGAGTGT 694

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RESULT 2
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LOCUS
DEFINITION
Yv56c11.s1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
IMAGE:230132 3', mRNA sequence.
ACCESSION
H74303.1 GI:1047714
VERSION
H74303.1
KEYWORDS
EST.
SOURCE
Homo sapiens (human)
ORGANISM
Homo sapiens
Eukaryota; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
1 (bases 1 to 414)
Hillier, L., Lennon, G., Becker, M., Bonaldo, M.F., Chiapelli, B.,
Chasse, S., Dietrich, N., DuBuque, T., Favello, A., Gish, W., Hawkins,
M., Hultman, M., Kucaba, T., Lacy, M., Le, M., Le, N., Mardis, E., Moore,
B., Morris, M., Parsons, J., Prange, C., Rifkin, L., Rohlfing, T.,
Schellenberg, K., Soares, M.B., Tan, F., Thierry-Mieg, J., Trevaskis, E.,
Underwood, K., Wohlmann, P., Waterston, R., Wilson, R. and Marra, M.
Generation and analysis of 280,000 human expressed sequence tags
Genome Res. 6 (9), 807-828 (1996)
97044478
8889549
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu

```

```

Insert Size: 1114
High quality sequence stops: 313
Source: IMAGE Consortium, LBNL
This clone is available royalty-free through LBNL; contact the
IMAGE Consortium (info@image.lbnl.gov) for further information.
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High quality sequence stop: 313.
Location/Qualifiers
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/clone="IMAGE:230132"
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/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
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/note="Organ: Liver and Spleen; Vector: p7T3D (Pharmacia)
with a modified polylinker; Site_1: Pac I; Site_2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo(dT) primer
15' AACTGGAGAATAAATGAAGATCTTTTTTTTTTTT 3',
double-stranded cDNA was ligated to Eco RI adapters
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified p7T3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT 75 a 127 c 116 g 95 t 1 others
ORIGIN
Query Match 41.3%; Score 409.8; DB 14; Length 414;
Best Local Similarity 99.3%; Pred No. 2.6e-48;
Matches 411; Conservative 0; Mismatches 3; Indels 0; Gaps 0;
QY 556 ATGGCAGGTCCACCTGCGAGCAGACACCATGCGAGGAGCTGCCAATGTCCTTCC 615
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QY 616 TGCGAGCGCAGACATCGGACTGTTCTGTCGAGGCTGCAACACCGCATGTCGAGC 675
Db 354 TGCGAGCGCAGACATCGGACTGTTCTGTCGAGGCTGCAACACCGCATGTCGAGC 295
QY 676 ACAGCGCCCTCACAGTGTGTCGCCCGCAGGTGTGTGACCCAGAGTGGAGACTGGCAGGTC 735
Db 294 ACAGCGCCCTCACAGTGTGTCGCCCGCAGGTGTGTGACCCAGAGTGGAGACTGGCAGGTC 235
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QY 856 TGTAAGATGAACCCCTCCAGAGAGCCAAACAGCGCAGAGGACTGCGAGCCATCAGCGTCA 915
Db 114 TGTAAGATGAACCCCTCCAGAGAGCCAAACAGCGCAGAGGACTGCGAGCCATCAGCGTCA 55
QY 916 CTGTTCTGATTTGGAGTTTCATGCAAAAATGAGTGTGTTTTAGCTGCTCTTGCCAC 969
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RESULT 3
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LOCUS
DEFINITION
Yv40b02.s1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
IMAGE:245163 3', mRNA sequence.
ACCESSION
N54458
VERSION
N54458.1 GI:1195778
KEYWORDS
EST.
SOURCE
Homo sapiens (human)

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Best Local Similarity 92.0%; Pred. No. 3.3e-43; Matches 439; Conservative 0; Mismatches 29; Indels 9; Gaps 4;	
QY	12 GGAACTAGAGGTTCTCACTGCCCCAGAGAGGCGCTACACCCAGGAGGATGGGGCTC 71
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QY	72 CTTGGGCTGTTCTGTTGGCGGTGCTGGCTGCCAGCAGTTCTCCAAAGGCAGGAGAA 131
Db	64 CTTGGGCTGTTCTGTTGGCGGTGCTGGCTGCCAGCAGTTCTCCAAAGGCAGGAGAA 123
QY	132 GAAATTAACCCCTGCTGCTTCCATTCCTTCAAAAGTCTCGAAGTTTTCCTCCAAAGGCGGC 191
Db	124 GAAATTAACCCCTGCTGCTTCCATTCCTTCAAAAGTCTCGAAGTTTTCCTCCAAAGGCGGC 183
QY	192 TGGGTGCTCATAACTGCTGTGCAACCCAGCAGCCACACCGCCCTACCTATTCCTCTGT 251
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QY	252 GGAACCAAGATCAAGGTGGCCAGAGAGTGTGAGAGCCACAGCGCGCTCTCTTC 311
Db	244 GGAACCAAGATCAAGGTGGCCAGAGAGTGTGAGAGCCACAGCGCGCTCTCTTC 303
QY	312 AACCTCAAGTGCACACTCACTCCAGTCACAGCTGCT-----CACCTACTTTCGCGGCGC 367
Db	304 AACCTCAAGTGCACACTCAAGTCCAGTCCAGCTGCTTCACTTATTTCTGCCGCGGC 363
QY	368 GTCTCCACTCAGTGGTCCCATGT-GGACAGTGCAGGCT---ACAGATGCATCTGGGAGC 423
Db	364 TTCTCCACTCAGTGGTCCCATGTGGGACAGTGCAGGCTTACAGATGGCATTTGGGAGC 423
QY	424 TGTGGTCCAGGCGAGTG-TCTGAGCTCGGGCCAACTTCACTCTGAGGACAGAGGG 479
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YR25C07.r1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone	
IMAGE:206316 5', mRNA sequence.	
H58326	VERSION
H58326.1	GT:1011158
EST.	
Homo sapiens (human)	SOURCE
Homo sapiens	ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;	REFERENCE
Mammalia; Euthera; Primates; Catarrhini; Hominidae; Homo.	AUTHORS
1 (bases 1 to 418)	
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J., Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevisan, E., Waterston, R., Williamson, A., Wohlmann, P., and Wilson, R.	
The WashU-Merck EST Project	
Unpublished	
Contact: Wilson RK	
Washington University School of Medicine	
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108	
Tel: 314 286 1800	
Fax: 314 286 1810	
Email: est@watson.wustl.edu	
Insert Size: 912	
High quality sequence stops: 251	
Source: IMAGE Consortium, LNL	
This clone is available royalty-free through LNL; contact the	
IMAGE Consortium (info@image.lnl.gov) for further information.	
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Seq primer: M13RP1	
High quality sequence stop: 251.	
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FEATURES	
source	


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MEDLINE      PUBMED      COMMENT
97044478
889549
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 1114
High quality sequence stops: 308
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
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High quality sequence stops: 308.
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/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares fetal liver spleen INFLS"
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with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo [drr] primer
[5', AACTGCAGAACTAATAAGATCTTTTTTTTTTTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pTVT3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
126 a 149 c 157 g 130 t 20 others
BASE COUNT
ORIGIN

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QY	10	CAGGAAC TAGGAGGTTCTCACTGCCCGACGACGA-GGCCCTACACCCACCGAGGCATGGGG	68	
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QY	69	CTCCCTGGGCTGTTCTGCTTGGCGTGCTGGCTGCCAGCAGC--TTCTCAAGGACACGGGA	127	
Db	61	CTCCCTGGGCTGTTCTGCTTGGCGTGCTGGCTGCCAGCAGCTTCTCCAAAGGACACGGGA	120	
QY	128	GGAAGAAATACCCCTGTGGTCTCCATTCGCTACAAAGTCTGGAAAGTTTTCCTCCAAAGG	187	
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QY	188	CGCGTGGGTGCTCATAAACCTGTGTGCACCCACGACACCGCCCATCACCTATTTCCT	247	
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QY	248	CTGTGGAACCAAGAACATCAAGGTGGCCAAAGAGTGTGAAGACCCACGAGCGCGCTTC	307	
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QY	308	CTTCAACCTCAACGTCACACTCAAGTCCAGTCCAGTCCAGACCT-GCTCACTACTT--CTGCGG	364	
Db	301	CTTCAACCTCAACGTCACACTCAAGTCCAGTCCAGTCCAGACCTGGCTCACTTAATTATGGCGG	360	
QY	365	GGCGTCTCTCCACCTCAGGTGCCCATGTGG---ACAGTGCCAGGCTACAGATGCACCTGGGA	421	
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QY	422	GCTGT---GGTCAAGCCAGTGTCTGAGCT---GGGGCCAACTTCATCTCAGGACAG	475	

Db 421 AGGTTGTGGTTCACAGCCAGTGTNTGAGTTTCCGGGNCATTTTAAATTTTCAGGACAG 481

QY 476 AGGGCAGGCCCCAGGGTGCAGATC--ATCTGCAGCGCGTCTCGGGCAGCCCACTAT 532

Db 481 AGGGNAAGNCCAGGTTGGAGTGTATTTGCCAGNTTTTATAGGAGTCCCAATTAT 540

RESULT 7

N76568 468 bp mRNA linear EST 28-JAN-1999

LOCUS YV40b02.r1 Soares fetal liver spleen INFLS Homo sapiens CDNA clone

DEFINITION IMAGE:245163 5', mRNA sequence.

ACCESSION N76568

VERSION N76568.1 GI:1239146

KEYWORDS EST.

SOURCE Homo sapiens (human)

ORGANISM Homo sapiens

REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo. 1 (bases 1 to 468)

AUTHORS Hillier,L., Lennon,G., Becker,M., Bonaldo,M.F., Chiapelli,B., Chisoe,S., Dietrich,N., DuBoue,T., Favello,A., Gish,W., Hawkins,M., Hultman,M., Kucaba,T., Lacy,M., Le,M., Le,N., Mardis,E., Moore,B., Morris,M., Parsons,J., Prange,C., Rifkin,L., Rohlfing,T., Schellenberg,K., Soares,M.B., Tan,F., Thierry-Mieg,J., Trevaskis,B., Underwood,K., Wohlmann,P., Waterston,R., Wilson,R. and Marra,M.

TITLE Generation and analysis of 280,000 human expressed sequence tags

JOURNAL Genome Res. 6 (9), 807-828 (1996)

MEDLINE 97044478

PUBMED 8889549

COMMENT Contact: Wilson RK Washington University School of Medicine 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108 Tel: 314 286 1800 Fax: 314 286 1810 Email: est@watson.wustl.edu This clone is available royalty-free through LLNL; contact the IMAGE Consortium (info@image.llnl.gov) for further information. Seq primer: reverse ET High quality sequence stop: 380.

FEATURES

source Location/Qualifiers

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/notes="Organ: Liver and Spleen; Vector: p7T3D (Pharmacia) with a modified polylinker; Site 1: Pac I; Site 2: Eco RI; 1st strand cDNA was primed with a Pac I - oligo(dT) primer [5' ACTCGAAGATTAATAAGACTTTTTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Pac I and cloned into the Pac I and Eco RI sites of the modified p7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fatima Bonaldo."

BASE COUNT 105 a 156 c 126 g 80 t 1 others

ORIGIN

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Best Local Similarity 90.3%; Pred. No. 1.2e-38;

Matches 457; Conservative 0; Mismatches 5; Indels 44; Gaps 7;

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 Db 14 GCCCAACCTCAGACACGATGGCGTCTCTTCAGCTGCTCTTTGCCATGTGGCTGC 73
 QY 104 CAGCAGCTTCTCAAAGCAGCGGAGGAAGAAATTACCCCTGTGGTTCCTAATGCCTACAA 163
 Db 74 CTGTGGCTTCTCAGAGGAGCAGACAGAAGGCATCA-----CCATTGCCTACAA 121
 QY 164 AGTCCTGGAAGTTTTCCCAGAACGCCTGGGTGCTCATTAACCTGCTGTGTGCACCCCAGCC 223
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 QY 224 ACCACGCCCATCACCTATTCCTCTGTGGAAACCAAGAACATCAAGTGTGCCAAGAAGGT 283
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 QY 404 GCTACAGATGCAGTGGGAGCTGTGGTCCAAGCCAGTGTCTGAGCTCGGGCCCAACTTCA 463
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 QY 464 TCTGCGGACAGAGGGCAGGCCCCAGGGTGGAGATGATCTGCCAGCGTCTCTCGGCGAG 523
 Db 422 CCTACGCCATGCGGAGCTCGGGGCCCACTGTGGAGCTGTCTGTGCTGGCATCTCAGCAG 481
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 Db 482 CCCCCCATCACCTACCGTCTGTGGGGAATGTGGGCGCTGTCTTTCAGCAGCAAAAGGCC 541
 QY 584 ATGCCACAGGCGCTGCCAACTTCTCTTCCTGCGGAGCAGACATCGGACTGTTCTG 643
 Db 542 ACTTCATGGNAACCAAGCCAACTTCTCCCTCCCGCTGTCTCCACACCACTGGTTGGTTCA 601
 QY 644 GTGCCAGGCTGCAAAACAACGCCAATGTCCAGCACAGCGCCCTCACAGTGTGTCGCCAGG 703


```

CC RZPD Deutsches Ressourcenzentrum fuer Genomforschung GmbH
CC Heubnerweg 6, D-14059 Berlin, Germany
CC Tel: +49 30 32639 101
CC Fax: +49 30 32639 111
CC www.rzpd.de
CC This clone is available royalty-free from RZPD;
CC contact RZPD (clone@rzpd.de) for further information.
CC Seq primer: T7, Primer sequence: TAATACGACTCACTATAGG
XX
FH Key Location/Qualifiers
FH source
FH 1..528
FT /db_xref="taxon:10090"
FT /note="1st strand cDNA was primed with a Not I - oligo(dT)
FT primer [5' TGTTACCAATCTGAAGTGGAGCGCGCGGAAATTTT
FT TTTTTTT 3'], on equal amounts of mRNA from 2
FT 13.5dpc and 2.14.5dpc embryos [total RNA provided by Minoru
FT Ko, Wayne State Univ., from 2 ]; double-stranded cDNA was
FT ligated to EcoRI adaptors 5'-AATTCGACAGG-3' and
FT 5'-CTTCGCGCG-3' (Pharmacia), digested with NotI and cloned
FT into the NotI and EcoRI sites of the pT7T3D-PacI vector.
FT Library went through one round of normalization, and was
FT constructed by Bento Soares and M.Fatima Bonaldo."
FT /organism="Mus musculus"
FT /clone="TMA9998K041131"
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Query Match 26.7%; Score 265.2; DB 4; Length 528;
Best Local Similarity 71.6%; Pred. No. 4.4e-28;
Matches 348; Conservative 0; Mismatches 138; Indels 0; Gaps 0;

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42 TCACCATTCCTTACAAAGTCTGGAGTTTATCCCAAGCGCGAGGGTGCTTAACT 101
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208 GTGTGTGACCCAGCCAGCCAGCCCATCACTATTCCCTCTGTGGAACCAAGACATCA 267
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
102 GCGATGCCCTGAGGGGTCCAGCCCATCATCATCTCTCTCTGGTAGCCAGGTATCC 161
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268 AGTGGCCAAAGAGTGTGTGAAGACCCAGAGCGCGCTTCTCAACCTCAACGTCAAC 327
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162 TGTGTGCAAAAAGGTTGTGCATGACTCCGTGCGCGCTTCTCAACATCAATATACCA 221
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282 ATGGACCCAGCAGAGCTCCAGATGTACCAGAACTGTGGGCTAAGCCAGTGTCTCAGC 341
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448 TGGGGCCAACTTCACTCTGAGGACAGAGGGGAGGCCGCCAGGGTGGAGATGATTCGC 507
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342 TGCAGGCTGACTTCTGCTTCACTGCTGAGGAGTCTGGGGCCCTCTGTGGAGCTGTCT 401
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
508 AGGCGTCTTGGGCGAGCCCACTATCAACCAAGCTGTATCGGGAAGGATGGCAGGTCC 567
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
402 TGGCATCTCTGAGCAGCCGCCCATCACTACCGCTTGTGGGAATGTGGCGGTTC 461
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
568 AACTGAGCAGAGACCATGCCAGAGCAGCTGCCAACTTCTCTTCCTGCCAGGCAGA 627
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
462 TTGCACAGCAAAAGGCCACTTATGGAATAACAGCCAACTTCTCTCTCCGCTGTCC 521
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
628 CATCGG 633
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
522 CCACTG 527

RESULT 15

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H40263/c
LOCUS
DEFINITION
IMAGE:191756 3', mRNA sequence.
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 384)
Hillier, L., Clark, N., Dubuque, T., Elliston, K., Hawkins, M., Holman
, M., Hultman, M., Kucaba, T., Le, M., Lennon, G., Marra, M., Parsons, J.,
Rifkin, L., Rohlfing, T., Soares, M., Tan, F., Trevaskis, E., Waterston
, R., Williamson, A., Wohlmann, P. and Wilson, R.
The WashU-Merck EST Project
Unpublished
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 1020
High quality sequence stops: 297
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Seq primer: Promega -21ml3
High quality sequence stop: 297.
Location/Qualifiers
1..384
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:3761545"
/db_xref="taxon:9606"
/clone="IMAGE:191756"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares fetal liver spleen INFILS"
/note="Organ: Liver and Spleen; Vector: pT7T3D (Pharmacia)
with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo(dT) primer
[5' AACGTGGAAGATTAATTAAGATCTTTTCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pT7T3 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT 70 a 122 c 111 g 77 t 4 others
ORIGIN
Query Match 26.3%; Score 261.2; DB 14; Length 384;
Best Local Similarity 98.6%; Pred. No. 1.8e-27;
Matches 273; Conservative 0; Mismatches 3; Indels 1; Gaps 1;
QY 693 GTGCCCCAGGTGGTGA-CCAGAGATGGAGACTGGCAGGTCCTCCCTGGAGAGCCCAT 751
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
278 GTGGGCCAGGTGGTGGACCCAGAGATGGAGACTGGCAGGTCCTCCCTGGAGAGCCCAT 219
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
752 CTTTCCCTTGGCGTCTACAGAGCACCCGCGCTCTGAGTGAAGAGGATTTGGGGGTT 811
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
218 CTTTCCCTTGGCGTCTACAGAGCACCCGCGCTCTGAGTGAAGAGGATTTGGGGGTT 159
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
812 CAGGTAGGGATGGGAGGTCAGAGACCGAAGCAGCAGCCATGTTAGATGAACCGTC 871
Db | | | | | | | | | | | | | | | | | | | | | | | | | | | |
158 CAGGTAGGGATGGGAGGTCAGAGACCGAAGCAGCAGCCATGTTAGATGAACCGTC 99
QY | | | | | | | | | | | | | | | | | | | | | | | | | | | |
872 CAGAGACCAAGCAGCGCAGAGGACTGCAGGCCATCAGGTGCACTGTTCTATTGGAG 931

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Db      98  CAGAGAGCAGCAGCGGACGAGGACTGCAGGCCATCAGCGTGCACCTGTCGTATTTGGAG 39
Qy      932  TTCTATGCAAAATGAGTGTTGTTTACGTCTTGCCA 968
Db      38  TTCTATGCAAAATGAGTGTTGTTTACGTCTTGCCA 2

RESULT 16
H58715/c
LOCUS      284 bp      mRNA      linear      EST 05-OCT-1995
DEFINITION Yr25c07.s1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
IMAGE:206316 3', mRNA sequence.
ACCESSION  H58715
VERSION     H58715.1 GI:1011547
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 284)
AUTHORS   Hallier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohlmann,P. and Wilson,R.
TITLE     The WashU-Merck EST Project
JOURNAL   Unpublished
COMMENT   Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wuston.wustl.edu
Insert Size: 912
High quality sequence stops: 241
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 912 Std Error: 0.00
Seq primer: Promega -21m3
High quality sequence stop: 241.
FEATURES             source
    1..284
        /organism="Homo sapiens"
        /mol_type="mRNA"
        /db_xref="GDB:3775447"
        /db_xref="taxon:9606"
        /clone="IMAGE:206316"
        /sex="male"
        /dev_stage="20 week-post conception fetus"
        /lab_host="DH10B (ampicillin resistant)"
        /clone_lib="Soares fetal liver spleen INFLS"
        /note="Organ: Liver and Spleen; Vector: pVT73D (Pharmacia)
with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo(dT) primer
[5', AACTGGAAGAATTAATAAGATCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pVT73 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT      50 a 100 c 68 g 66 t
ORIGIN
Query Match      25.9%; Score 256.8; DB 14; Length 284;
Best Local Similarity 99.2%; Pred. No. 8.7e-27;
Matches 258; Conservative 0; Mismatches 2; Indels 0; Gaps 0;

Qy      693  GTGCCCCAGTGGTGCACAGAGATGGAGGATGCGAGGTCCTCGAGAGCCCATC 752
Db      260  GTGGGCCAGTGGTGCACAGAGATGGAGGATGCGAGGTCCTCGAGAGCCCATC 201
Qy      753  CTTGCCTTCCGCTCTACAGGAGCACCGCGCTCTGAGTGAAGAGGAGTTTGGGGGTTTC 812

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Db      200  CTTGCCTTCCGCTCTACAGGAGCACCGCGCTCTGAGTGAAGAGGAGTTTGGGGGTTTC 141
Qy      813  AGGATAGGGAATGGGGAGGTCAGAGGACGCAAGCAGCAGCCATGTAGAAATGAACCGTCC 872
Db      140  AGGATAGGGAATGGGGAGGTCAGAGGACGCAAGCAGCAGCCATGTAGAAATGAACCGTCC 81
Qy      873  AGAGAGCAAGCAGCGCAGAGGACTGCAGGCCATCAGCGTGCACCTGTCGTATTTGGAGT 932
Db      80  AGAGAGCAAGCAGCGCAGAGGACTGCAGGCCATCAGCGTGCACCTGTCGTATTTGGAGT 21
Qy      933  TCATGCAAAATGAGTGTTGTTT 952
Db      20  TCATGCAAAATGAGTGTTGTTT 1

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```

RESULT 17
T80187/c
LOCUS      376 bp      mRNA      linear      EST 15-MAR-1995
DEFINITION Yd8593.s1 Soares fetal liver spleen INFLS Homo sapiens cDNA clone
IMAGE:115060 3', mRNA sequence.
ACCESSION  T80187
VERSION     T80187.1 GI:698696
KEYWORDS   EST.
SOURCE     Homo sapiens (human)
ORGANISM   Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE  1 (bases 1 to 376)
AUTHORS   Hillier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman
,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J.,
Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaskis,E., Waterston
,R., Williamson,A., Wohlmann,P. and Wilson,R.
TITLE     The WashU-Merck EST Project
JOURNAL   Unpublished
COMMENT   Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@wuston.wustl.edu
Insert Size: 1048
High quality sequence stops: 330
Source: IMAGE Consortium, LLNL
This clone is available royalty-free through LLNL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1048 Std Error: 0.00
Seq primer: -21m13
High quality sequence stop: 330.
FEATURES             Location/Qualifiers
    1..376
        /organism="Homo sapiens"
        /mol_type="mRNA"
        /db_xref="GDB:470677"
        /db_xref="taxon:9606"
        /clone="IMAGE:115060"
        /sex="male"
        /dev_stage="20 week-post conception fetus"
        /lab_host="DH10B (ampicillin resistant)"
        /clone_lib="Soares fetal liver spleen INFLS"
        /note="Organ: Liver and Spleen; Vector: pVT73D (Pharmacia)
with a modified polylinker; Site 1: Pac I; Site 2: Eco RI;
1st strand cDNA was primed with a Pac I - oligo(dT) primer
[5', AACTGGAAGAATTAATAAGATCTTTTCTTTTCTTTT 3'],
double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Pac I and cloned into the Pac I
and Eco RI sites of the modified pVT73 vector. Library
went through one round of normalization. Library
constructed by Bento Soares and M.Fatima Bonaldo."
BASE COUNT      65 a 117 c 114 g 78 t 2 others
ORIGIN
Query Match      23.7%; Score 235.2; DB 14; Length 376;
Best Local Similarity 94.4%; Pred. No. 8.2e-24;
Matches 254; Conservative 0; Mismatches 14; Indels 1; Gaps 1;

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Db 400 TG 401

RESULT 23
T87491

LOCUS
DEFINITION
T87491.1 GI:715843
EST.
KEYWORDS
SOURCE
ORGANISM
Homo sapiens (human)
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
REFERENCE
AUTHORS
1 (bases 1 to 324)
Hallier,L., Clark,N., Dubuque,T., Elliston,K., Hawkins,M., Holman,M., Hultman,M., Kucaba,T., Le,M., Lennon,G., Marra,M., Parsons,J., Rifkin,L., Rohlfing,T., Soares,M., Tan,F., Trevaekis,E., Waterston,R., Williamson,A., Wohldmann,P. and Wilson,R.
TITLE
The WashU-Merck EST Project
JOURNAL
Unpublished
COMMENT
Contact: Wilson RK
Washington University School of Medicine
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: est@watson.wustl.edu
Insert Size: 1048
High quality sequence stops: 179 Source: IMAGE Consortium, LMLL
This clone is available royalty-free through LMLL ; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
Insert Length: 1048 Std Error: 0.00
Seq primer: M13RP1
High quality sequence stop: 179.

FEATURES
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1..324
Location/Qualifiers
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="GDB:470677"
/db_xref="taxon:9606"
/clone="IMAGE:115060"
/sex="male"
/dev_stage="20 week-post conception fetus"
/lab_host="DH10B (ampicillin resistant)"
/clone_lib="Soares fetal liver spleen lNFTS"
/note="Organ: Liver and Spleen; Vector: pT7T3D (Pharmacia) with a modified polylinker; Site_1: Pac I; Site_2: Eco RI; 1st strand cDNA was primed with a Pac I - oligo(dT) primer [5'- AACTGAGAAATTAATTAAGATCTTTTTTTTTTTTTTTT 3'], double-stranded cDNA was ligated to Eco RI adaptors (Pharmacia), digested with Pac I and cloned into the Pac I and Eco RI sites of the modified pT7T3 vector. Library went through one round of normalization. Library constructed by Bento Soares and M.Fatima Bonaldo."

BASE COUNT
ORIGIN
70 a 99 c 77 g 78 t

Query Match 19.4%; Score 192.2; DB 14; Length 324;
Best local Similarity 97.7%; Pred. No. 9.4e-18;
Matches 216; Conservative 0; Mismatches 3; Indels 2; Gaps 2;

QY 131 AGAAATTACCCCTGTGCTCTCCATTGCTACAAAGTCCTGGAAGTTTCCCAAGGCCG 190
|||||
Db 75 AGAAATTACCCCTGTGCTCTCCATTGCTACAAAGTCCTGGAAGTTTCCCAAGGCCG 134
|||||

QY 191 CTGGTGCTCATTAACCTGCTGTGCACCCAGCCACACCGCCCATACCTATTTCCTCTG 250
|||||

Db 135 CTGGTGCTCATTAACCTGCTGTGCACCCAGCCACACCGCCCATACCTATTTCCTCTG 194
|||||

QY 251 TGGAAACAAGAAATCAAGT-TGGCAAGAGTGGTGAAGCCACGAGCGCGCTCC- 308
|||||

Db 195 TGGAAACAAGAAATCAAGTGGCGCAAGAGTTGTGAAGCCACGAGCGCGCTCT 254
|||||

QY 309 TTCAACTCAAGTTCACACTCAAGTTCAGTCCAGACCTGCT 349
 Db 255 TTCAACTTAACTTCACACTCAAGTCCAGTCCAGACCTGGT 295

RESULT 24
 W65713
 LOCUS 467 bp mRNA linear EST 11-JUN-1996
 DEFINITION me0906.r1 Soares mouse embryo NbME13.5 14.5 Mus musculus cDNA
 clone IMAGE:387034 5', mRNA sequence.
 ACCESSION W65713
 VERSION W65713.1 GI:1373922
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 (Bases 1 to 467)
 AUTHORS Marra, M., Hillier, L., Allen, M., Bowles, M., Dietrich, N., Dubuque, T.,
 Geisel, S., Kucaba, T., Lacy, M., Le, M., Martin, J., Morris, M.,
 Schellenberg, K., Stepcoe, M., Tan, F., Underwood, K., Moore, B.,
 Theising, B., Wylie, T., Lennon, G., Soares, B., Wilson, R. and
 Waterston, R.
 TITLE The WashU-HMI Mouse EST Project
 JOURNAL Unpublished
 COMMENT Contact: Marra M/Mouse EST Project
 WashU-HMI Mouse EST Project
 Washington University School of MedicineP
 4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
 Tel: 314 286 1800
 Fax: 314 286 1810
 Email: mouseest@watson.wustl.edu
 This clone is available royalty-free through LML; contact the
 IMAGE Consortium (info@image.llnl.gov) for further information.
 MG1:238866
 Seq primer: ETPPrimer
 High quality sequence stop: 345.
 Location/Qualifiers
 1. .467
 /organism="Mus musculus"
 /mol_type="mRNA"
 /strain="C57BL/6J"
 /db_xref="taxon:10090"
 /clone="IMAGE:387034"
 /sex="unknown"
 /tissue_type="embryo"
 /dev_stage="13.5-14.5dpc total fetus"
 /lab_host="DH10B"
 /clone_lib="Soares mouse embryo NbME13.5 14.5"
 /notes="Vector: pT7T3D-Pac (Pharmacia) with a modified
 polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
 was primed with a Not I - oligo(dT) primer [5',
 TGTTCACATCTGAAGTGGAGCGCGCGGAATTTTGTGTGTGTGTGTGT
 T 3'], on equal amounts of mRNA from 2 13.5dpc and 2
 14.5dpc embryos [total RNA provided by Minoru Ko, Wayne
 State Univ., from 2]; double-stranded cDNA was ligated to
 Eco RI adaptors (Pharmacia), digested with Not I and
 cloned into the Not I and Eco RI sites of the modified
 pT7T3 vector. Library went through one round of
 normalization, and was constructed by Bento Soares and
 M.Fatima Bonaldo."
 BASE COUNT 97 a 159 c 112 g 99 t
 ORIGIN

Query Match 18.4%; Score 182.2; DB 14; Length 467;
 Best Local Similarity 68.0%; Pred.No.2e-16;
 Matches 319; Conservative 0; Mismatches 133; Indels 17; Gaps 4;

QY 66 GGCGTCCCTGGGGTGTTGTGCTGGCGTGGTCCAGCAGCTTCTCCAGGACGG 125
 Db 14 GGCGTCTTCAGCTGCTCTCTTTGGCATGCTGGCTGCTGGTCTCAGAGGACGAG 73

Murata, M., Nakamura, M., Nomura, K., Numazaki, R., Ohno, M., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shiraki, T., Tagami, M., Waki, K., Watahiki, A., Muramatsu, M. and Hayaishizaki, Y. Direct Submission

Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome 12, 673-677 (2001)

Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multicapillary sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN, Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Dr. John Todd (Dept. of Medical Genetics Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome Trust/MRC building Addenbrookes Hospital Cambridge) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES
source

Location/Qualifiers
1. .364
/organism="Mus musculus"
/mol_type="mRNA"
/strain="M0D"
/db_xref="taxon:10090"
/clone="F830016f14"
/tissue_type="activated spleen"
/clone_lib="RIKEN full-length enriched, activated spleen"
84 a 123 c 82 g 74 t 1 others

Query Match 17.6%; Score 174.6; DB 13; Length 364;
Best Local Similarity 69.6%; Pred. No. 2.6e-15;
Matches 256; Conservative 0; Mismatches 100; Indels 12; Gaps 1;
QY 44 GCCTACACCCACGAGGATGGGCTCCCTGGCTGTTCTGCTGGCCGTGCTGCTGC 103
Db 9 GCCCCAACCTCAGAGGATGGGCTCCCTTCAGCTGCTCTTTGGCAATGCTGCTGC 68
QY 104 CAGCAGTTCTCAAGCAGCGGAGGAGAAATACCCCTGTGGTCTCCATTGCCCTACAA 163
Db 69 CTGTGGCTTCTCAGAGGAGCAGAGAAGGCATCA-----CCATTGCCCTACAA 116
QY 164 AGTCCTGGAGTTTCCCAAGCCGCTGGTGTCTATACCTGTGTCACCCCGAGCC 223
Db 117 AGTACTGGAAGTTTATCCCAAGCCGAGGGGTGCTTATAACTGGGATGCCCTGAGGC 176
QY 224 ACCACGCCCATCACCTATTCCCTGTGGAAACCAAGAACATCAAGTGGCCCAAGAGGT 283
Db 177 GTCCAGCCATACATATCTCTCTGGCTAGCCGAGGTATCTGGTGGCAAAAAGGT 236
QY 284 GGTGAAGACCCAGCGCCGCTCTTCAACCTCAAGCTCACACTCAAGTCCAGTCCAGA 343
Db 237 TGTGCAATGATCCGTGCGCGCTCTTCAACATCATATCACCATCAAGTCCAGGCCAGA 296
QY 344 CTGTCTACCTACTCTGCGGGGGTCTCTCACTCAGGTGCCCATGTGACAGTCCAG 403
Db 297 CCGTCTACCTACTCTGCGCAGGCACTCGAATCTGGCACCTATGGACCCAGCAGNAG 356
QY 404 GCTACAGA 411
Db 357 GTTCCAGA 364

RESULT 27
BI347002

LOCUS
DEFINITION
ACCESSION
VERSION
KEYWORDS
SOURCE
ORGANISM

REFERENCE
AUTHORS

TITLE

JOURNAL
MEDLINE
PUBMED
COMMENT

FEATURES
source

Location/Qualifiers
1. .537
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH108"
/clone_lib="MARC 2P1G"
/note="Vector: pCMV SPOT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, ovary,
endometrium, hypothalamus, pituitary, and placenta."
110 a 182 c 154 g 91 t

BASE COUNT
ORIGIN

Query Match 15.5%; Score 154; DB 12; Length 537;
Best Local Similarity 75.2%; Pred. No. 1.7e-12;
Matches 206; Conservative 0; Mismatches 65; Indels 3; Gaps 1;
QY 433 AGCCAGTGTCTGAGCTGCGGGCCCACTTCATCTGTGAGGACAGAGG---GGCAGGCCCA 489
Db 132 AGCCCGTGTCCAGCTGTGGTTAACTTCATCTGTGACGAGCCCGTGGGGCCCT 191
QY 490 GGTGTGAGATGATCTGCCAGGCGTCTCGGGCAGCCACCTATCACCAAGCCTGATCG 549
Db 192 GGTGTGAGTGTCTGTCTGTGGCATCTCTGGGAGCCCACTTACCTAGTCTTGTGCG 251
QY 550 GGAAGGATGGCAGGTCCACCTGTGACGAGACCATGCCCACAGGCGCTGCCAACTTCT 609
Db 252 GGAGGGACAGTGGCACCTTACATGAGGACACACAGAACTACAAGGAGTGTCCACTTCT 311
QY 610 CTTTCTGCGGAGCCAGACATCGATCTGTGTTCTGTGCGAGGCTGCAACACAGCCAAATG 669
Db 312 CTTTCCCGTCACTCCAGAGCTGTGTGTGGTCCAGGTCCAGGCTGCAACACATCAGCG 371
QY 670 TCCAGCAGACGCCCTCAGTGTGTCGCCCCAGG 703
Db 372 CCAGCAGAGTGCCTTCACATGGTGGCCCCCAGG 405

RESULT 28
BI347003

BI347002 537 bp mRNA linear EST 30-JUL-2001
376394 MARC 2P1G Sus scrofa cDNA 5', mRNA sequence.
BI347002
EST.
BI347002.1 GI:15040291
KEYWORDS
Sus scrofa (pig)
Sus scrofa
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
1 (Bases 1 to 537)
Fahrenkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J.,
Vallat, J., Wise, T., Rohrer, G.A., Perte, G., Sultana, R., Quackenbush,
J. and Keeler, J.W.
Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
Mamm. Genome 13 (8), 475-478 (2002)
22213789
12226715

Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@mail.marc.usda.gov
Single pass sequencing. Bases called and alt. trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -mismatch 12 options.
PCR Primers

FORWARD: AGGAAACAGCTATGACCAT
BACKWARD: GTTTCCAGTCCAGCAGC
Plate: 130 row: N column: 7
Seq primer: ATTAGTGCACACTATAG.

LOCUS BI347003 376395 MARC 2P1G Sus scrofa cDNA 5', mRNA linear EST 30-JUL-2001
 DEFINITION 376395 MARC 2P1G Sus scrofa cDNA 5', mRNA sequence.
 ACCESSION BI347003
 VERSION BI347003.1 GI:15040292
 KEYWORDS EST.
 SOURCE Sus scrofa (pig)
 ORGANISM Sus scrofa
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
 REFERENCE 1 (bases 1 to 582)
 AUTHORS Fahrénkrug, S.C., Smith, T.P.L., Freking, B.A., Cho, J., White, J.,
 Vallet, J., Wise, T., Rohrer, G.A., Perce, G., Sultana, R., Quackenbush,
 J., and Keele, J.W.
 TITLE Porcine gene discovery by normalized cDNA-library sequencing and
 EST cluster assembly
 JOURNAL Mamm. Genome 13 (8), 475-478 (2002)
 MEDLINE 22213789
 PUBMED 12226715
 COMMENT Contact: Smith TPL
 USDA, ARS, US Meat Animal Research Center
 PO Box 166, Clay Center, NE 68933-0166, USA
 Tel: 402 762 4366
 Fax: 402 762 4390
 Email: smith@email.marc.usda.gov
 Single pass sequencing. Bases called and alt_trimmed with phred
 v0.980904.e. Vector identified by cross_match with the -minscore 18
 and -minmatch 12 options.
 PCR Primers
 FORWARD: AGGAACAGCTATGACCAT
 BACKWARD: GTTTCACGTCAGGCG
 Plate: 130 row: N column: 8
 Seq primer: ATTATGGTGACACTATAG.
 Location/Qualifiers
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 /organism="Sus scrofa"
 /mol_type="mRNA"
 /db_xref="taxon:9823"
 /tissue_type="pooled"
 /lab_host="DH10B"
 /clone_lib="MARC 2P1G"
 /note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
 Library made from pooled tissue from testis, ovary,
 endometrium, hypothalamus, pituitary, and placenta."

FEATURES

source

BASE COUNT 122 a 191 c 166 g 103 t
 ORIGIN
 Query Match 15.5%; Score 154; DB 12; Length 582;
 Best Local Similarity 75.2%; Pred. No. 1.7e-12;
 Matches 206; Conservative 0; Mismatches 65; Indels 3; Gaps 1;
 QY 433 AGCCAGTGTCTGAGTGGGCGCACTTCACTCTGCAGGACAGAGG---GGCAGGCCCA 489
 Db 132 AGCCGCTGCCAGTGTGTTTAACTTTCATCTCTGCGAGGCGCCGTCGGGCCCTT 191
 QY 490 GGGTGGAGATGATCTGCGAGCGTCTCTGGGAGCCACCTATCACCAACAGCCTGATCG 549
 Db 192 GGGTGAAGGTGTGCTGTGGGATCTCTGGGAGCCACCTATCACCTACAGTCTTGTGCG 251
 QY 550 GGAAGGATGGGAGGTCTCCACTTCGAGCAGAGACCATGCCAGGCGCTGCCAATCTCT 609
 Db 252 GGAGGGACAGTGGCGACCTTACATGAAGCAGACACAGAACTACAGGAAGCTGCCAATCTCT 311
 QY 610 CTTCTCTCGAGGCAGACATCGGACTGGTCTGTGTGTCAGGCTGCAACACAGCCCAATG 669
 Db 312 CTTCTCCGCTCACCAGACGCTGCTGCTGGCTCAGGTGCGAGCTGCAACACCAATCAGCG 371
 QY 670 TCCAGCAGCGCCCTCACTGTGTCGCCCCAGG 703
 Db 372 CCCAGCAGTGGCCCTCACTGTGTCGCCCCAGG 405
 RESULT 29
 CH051335

LOCUS CB051335 443 bp mRNA linear EST 17-JAN-2003
 DEFINITION NISC gj23h11.y1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:3289604
 5', mRNA sequence.
 ACCESSION CB051335
 VERSION CB051335.1 GI:27789622
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Homo sapiens
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
 Mammalia; Eutheria; Primates; Catarrhini; Homnidae; Homo.
 REFERENCE 1 (bases 1 to 443)
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
 Tumor Gene Index
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaaps-remail.nih.gov
 CDNA Library Preparation:
 CDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
 DNA Sequencing by: National Institutes of Health Intramural
 Sequencing Center (NISC)
 Clone distribution: NCI-CGAP clone distribution information can be
 found through the I.M.A.G.E. Consortium/LLNL at:
 info@image.llnl.gov
 Plate: L1AM8055 row: P column: 21
 Seq primer: M13RP1 reverse primer (ABI).
 Location/Qualifiers
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 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:3289604"
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 /dev_stage="adult"
 /lab_host="DH10B"
 /clone_lib="NCI_CGAP_Pr28"
 /note="Organ: prostate; Vector: pT7T3D-Pac (Pharmacia)
 with a modified polylinker; Plasmid DNA from the
 normalized library NCI_CGAP_Pr22 was prepared, and ss
 circles were made in vitro. Following HAP purification,
 this DNA was used as tracer in a subtractive hybridization
 reaction. The driver was PCR-amplified cDNAs from a pool
 of 5,000 clones made from the same library (cloneIDs
 985608-986759, 1101192-1101959, and 1217928-1220615).
 Subtraction by Bento Soares and M. Fatima Bonaldo."

BASE COUNT 129 a 112 c 111 g 91 t
 ORIGIN
 Query Match 15.5%; Score 153.8; DB 14; Length 443;
 Best Local Similarity 98.7%; Pred. No. 2e-12;
 Matches 155; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 833 CAGAGGACGCAAGACGACGCGCATGTAGATGAACCTCCAGAGAGCAAGCAGCGAGA 892
 Db 250 CAGAGGACGCAAGACGACGCGCATGTAGATGAACCTCCAGAGAGCAAGCAGCGAGA 309
 QY 893 GGACTCGAGCCCATCAGCGTGCACCTGTTCGTATTGGAGTTTCATGCAAAATGAGTGTGT 952
 Db 310 GGACTCGAGCCCATCAGCATGCATCTTCGTATTGGAGTTTCATGCAAAATGAGTGTGT 369
 QY 953 TTAGCTGCTTTCGCCACAAAAAATAAAAAAAAAAAAAA 989
 Db 370 TTAGCTGCTTTCGCCACAGAAAAAATAAAAAAAAAAAAAA 406
 RESULT 30
 CB051334/c
 LOCUS CB051334 540 bp mRNA linear EST 17-JAN-2003
 DEFINITION NISC gj23h11.x1 NCI_CGAP_Pr28 Homo sapiens cDNA clone IMAGE:3289604
 3', mRNA sequence.
 ACCESSION CH051334
 VERSION CB051334.1 GI:27789621
 KEYWORDS EST.

SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 540)
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-r@mail.nih.gov
 cDNA Library Preparation:
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
 DNA Sequencing by: National Institutes of Health Intramural Sequencing Center (NISC)
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
 Plate: LLAM8055 row: P column: 21
 Seq primer: -21M13 forward primer (ABI).
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 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:3289604"
 /sex="male"
 /dev_stage="adult"
 /lab_host="DH10B"
 /notes="Organ: prostate; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Plasmid DNA from the normalized library NCI-CGAP Pr22 was prepared, and ss this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonoids 985608-986759, 1101192-1101959, and 1217928-1220615). Subtraction by Bento Soares and M. Fatima Bonaldo."
 BASE COUNT 126 a 113 c 128 g 173 t
 ORIGIN
 Query Match 15.5%; Score 153.8; DB 14; Length 540;
 Best Local Similarity 98.7%; Pred. No. 1.8e-12;
 Matches 155; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
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 Db 305 CAGAGCAGCAAGCAGCAGCCATGTAGAAATGAACCGTCCAGAGAGCCCAAGCAGCAGA 246
 QY 893 GGACTCAGGCCATCAGCGTGCACCTGTTGTTATTTGGAGTTCATGCAAAATGAGTGTGT 952
 Db 245 GGACTCAGGCCATCAGCGTGCACCTGTTGTTATTTGGAGTTCATGCAAAATGAGTGTGT 186
 QY 953 TTAGCTGCTCTTCCCAAAAAAATAAAAAAAAAAAAAA 969
 Db 185 TTAGCTGCTCTTCCCAAAAAAATAAAAAAAAAAAAAA 149
 RESULT 31
 LOCUS N88474
 DEFINITION X3475F Human fetal heart, Lambda ZAP Express Homo sapiens cDNA clone K3475 5', mRNA sequence.
 ACCESSION N88474
 VERSION N88474.1 GI:1441676
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 194)
 AUTHORS Liew, C.C.

SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 540)
 AUTHORS NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
 TITLE National Cancer Institute, Cancer Genome Anatomy Project (CGAP), Tumor Gene Index
 JOURNAL Unpublished
 COMMENT Contact: Robert Strausberg, Ph.D.
 Email: cgaps-r@mail.nih.gov
 cDNA Library Preparation:
 cDNA Library Arrayed by: The I.M.A.G.E. Consortium/LLNL
 DNA Sequencing by: National Institutes of Health Intramural Sequencing Center (NISC)
 Clone distribution: NCI-CGAP clone distribution information can be found through the I.M.A.G.E. Consortium/LLNL at: info@image.llnl.gov
 Plate: LLAM8055 row: P column: 21
 Seq primer: -21M13 forward primer (ABI).
 FEATURES
 source
 1..540
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
 /clone="IMAGE:3289604"
 /sex="male"
 /dev_stage="adult"
 /lab_host="DH10B"
 /notes="Organ: prostate; Vector: pT7T3D-Pac (Pharmacia) with a modified polylinker; Plasmid DNA from the normalized library NCI-CGAP Pr22 was prepared, and ss this DNA was used as tracer in a subtractive hybridization reaction. The driver was PCR-amplified cDNAs from a pool of 5,000 clones made from the same library (clonoids 985608-986759, 1101192-1101959, and 1217928-1220615). Subtraction by Bento Soares and M. Fatima Bonaldo."
 BASE COUNT 126 a 113 c 128 g 173 t
 ORIGIN
 Query Match 15.5%; Score 153.8; DB 14; Length 540;
 Best Local Similarity 98.7%; Pred. No. 1.8e-12;
 Matches 155; Conservative 0; Mismatches 2; Indels 0; Gaps 0;
 QY 833 CAGAGCAGCAAGCAGCAGCCATGTAGAAATGAACCGTCCAGAGAGCCCAAGCAGCAGA 892
 Db 305 CAGAGCAGCAAGCAGCAGCCATGTAGAAATGAACCGTCCAGAGAGCCCAAGCAGCAGA 246
 QY 893 GGACTCAGGCCATCAGCGTGCACCTGTTGTTATTTGGAGTTCATGCAAAATGAGTGTGT 952
 Db 245 GGACTCAGGCCATCAGCGTGCACCTGTTGTTATTTGGAGTTCATGCAAAATGAGTGTGT 186
 QY 953 TTAGCTGCTCTTCCCAAAAAAATAAAAAAAAAAAAAA 969
 Db 185 TTAGCTGCTCTTCCCAAAAAAATAAAAAAAAAAAAAA 149
 RESULT 31
 LOCUS N88474
 DEFINITION X3475F Human fetal heart, Lambda ZAP Express Homo sapiens cDNA clone K3475 5', mRNA sequence.
 ACCESSION N88474
 VERSION N88474.1 GI:1441676
 KEYWORDS EST.
 SOURCE Homo sapiens (human)
 ORGANISM Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
 REFERENCE 1 (bases 1 to 194)
 AUTHORS Liew, C.C.

cdNAs from fetal heart (1996)
 Unpublished
 Contact: Liew CC
 Brigham and Women's Hospital
 Harvard Medical School
 75 Francis St. Boston, MA 02115, USA
 Tel: 6177328915
 Fax: 6179750995
 Email: cliu@rics.bwh.harvard.edu
 Seq primer: GAAATTAACCTCCTCACTAAAGG.
 FEATURES
 Location/Qualifiers
 1..194
 /organism="Homo sapiens"
 /mol_type="mRNA"
 /db_xref="taxon:9606"
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 /clone_lib="Human fetal heart, Lambda ZAP Express"
 /note="Vector: Lambda ZAP Express; Site 1: EcoRI; Site 2: XhoI; mRNA was purified from human fetal hearts (8-10 weeks). cDNA was synthesized using a XhoI-Oligo dT adaptor-primer. EcoRI adaptors were ligated, followed by digestion with XhoI, for directional cloning into predigested lambda ZAP Express."
 BASE COUNT 51 a 43 c 63 g 37 t
 ORIGIN
 Query Match 14.6%; Score 145.2; DB 14; Length 194;
 Best Local Similarity 96.5%; Pred. No. 4.6e-11;
 Matches 191; Conservative 0; Mismatches 3; Indels 4; Gaps 4;
 QY 750 ATCCCTGCTCCGCTCTACAGGAGCACCCTGCTGAGTGAAGAGAGTTGGGGG 809
 Db 1 ATCCCTGCTCCGCTCTACAGGAGCACCCTGCTGAGTGAAGAGAGTTGGGGG 59
 QY 810 TTCAGATAGGAATGGGAGGTTCAGAGGACCAAGCAGCAGCCATGTAGATGAACG 869
 Db 60 TTCAGATAGGAATGGGAGGTTCAGAGGACCAAGCAGCAGCCATGTAGATGAACG 119
 QY 870 TCCAGAGAGCCCAAGCAGCAGGAGTTCAGGAGCCTCAGGCGCATCGCTGCTGATTTGG 929
 Db 120 T-CAGAGAGCCCAAGCAGCAGGAGTTCAGGAGCCTCAGGCGCATCGCTGCTGATTTGG 176
 QY 930 AGTTCATGCAAAATGAGT 947
 Db 177 AGTTCATGCAAAATGAGT 194
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 LOCUS BB842514
 DEFINITION BB842514 RIKEN full-length enriched, 6 days neonate spleen Mus musculus cDNA clone F430005P21 5', mRNA sequence.
 ACCESSION BB842514
 VERSION BB842514.1 GI:17043245
 KEYWORDS EST.
 SOURCE Mus musculus (house mouse)
 ORGANISM Mus musculus
 Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi; Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
 REFERENCE 1 (bases 1 to 382)
 AUTHORS Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T., Hayatsu, N., Hiramoto, K., Hiraoka, T., Hirozane, T., Imotani, K., Ishii, Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T., Nakamura, M., Nishi, K., Nomura, K., Numasaki, R., Okazaki, Y., Okido, T., Saito, R., Sakai, C., Sakai, K., Sakazume, N., Sasaki, D., Sato, K., Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa, A., Takahashi, F., Takaku-Akahira, S., Tanaka, T., Tomaru, A., Toya, T., Watabiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.
 RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al. 2001)
 JOURNAL Unpublished
 COMMENT Contact: Yoshihide Hayashizaki

Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216

Email: genome-res@gsr.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh
M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.
Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura
S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and
Hayashizaki, Y.
RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara
Y. and Hayashizaki, Y.

Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.
e mouse tissues.

FEATURES Location/Qualifiers
Source
1..382

/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="F43005P21"
/tissue_type="spleen"
/dev_stage="6 days neonate"
/clone_lib="RIKEN full-length enriched, 6 days neonate
spleen"

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ORIGIN
Query Match 14.4%; Score 143; DB 10; Length 382;
Best Local Similarity 69.8%; Pred. No. 7e-11;
Matches 213; Conservative 0; Mismatches 80; Indels 12; Gaps 1;

QY 44 GCCCTACACCCAGGAGGATGGGGCTCCCTGGGCTGTCTGTGGCGTGGTGGCGTGGCTGC 103
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QY 104 CAGCAGCTTCTCAAGGACGGGAGGAAGAAATPACCCCTGTGCTCTCCATTGCTACAA 163
DB 150 CTGTGGCTTCTCAGAGGAGCAGACAGAGGCATCA-----CCATTGCTACAA 197
QY 164 AGTCTCGAGTCTTCCCAAGGCGCTGGGTCTCATACCTGCTGTGCAACCCAGCC 223
DB 198 AGTACTGGAAGTTATCCCAAGCCGGAGGGTCTTTAATACCTGCGATGCCCTGAGGC 257
QY 224 ACCACCCGCCCATCACCTATTCCCTCTGTGGAACCAAGAACATCAAGTGGCCAGAGGT 283
DB 258 GTCCAGCCCATCATACACTCTCTCTGGGTAGCGAGGTATCTGTGGTGAAGAAAGGT 317
QY 284 GGTGAACCCACGAGCGGCGCTCTTCAACCTCAACGTCACACTCAAGTCCAGTCCAGA 343
DB 318 TGTGCATGACTCCGTGGCGGCTCTTCAACATCATCAATATCAGCATCAAGTCCAGCCAGA 377
QY 344 CCTGC 348
DB 378 CCTGC 382

RESULT 33
BB843270

LOCUS BB843270 370 bp mRNA linear EST 26-NOV-2001
DEFINITION BB843270 RIKEN full-length enriched, 6 days neonate spleen Mus
musculus cDNA clone F430010H21 5', mRNA sequence.
ACCSSION BB843270
VERSION BB843270.1 GI:17081637
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
REFERENCE 1 (bases 1 to 370)
AUTHORS Akimura, T., Arakawa, T., Carninci, P., Furuno, M., Hanagaki, T.,
Y., Ito, M., Kawai, J., Kojima, Y., Konno, H., Kouda, M., Matsuyama, T.,
Nakamura, M., Nishi, K., Numata, K., Numasaki, R., Okazaki, Y., Okado, T.,
Saito, R., Sakai, C., Sakai, K., Sakazume, N., Sasaki, D., Sato, K.,
Shibata, K., Shinagawa, A., Shiraki, T., Sogabe, Y., Suzuki, H., Tagawa
A., Takahashi, F., Takaku-Akahisa, S., Tanaka, T., Tomaru, A., Toya, T.,
Watahiki, A., Yasunishi, A., Muramatsu, M. and Hayashizaki, Y.
RIKEN Encyclopedia of Mouse Full-length cDNAs (Akimura, T., et al.
2001)

Unpublished
Contact: Yoshihide Hayashizaki
Laboratory for Genome Exploration Research Group, RIKEN Genomic
Sciences Center (GSC), Yokohama Institute
The Institute of Physical and Chemical Research (RIKEN)
1-7-22 Suehiro-cho, Tsurumi-ku, Yokohama, Kanagawa 230-0045, Japan
Tel: 81-45-503-9222
Fax: 81-45-503-9216
Email: genome-res@gsr.riken.go.jp,
URL: http://genome.gsc.riken.go.jp/
Carninci, P., Shibata, Y., Hayatsu, N., Sugahara, Y., Shibata, K., Itoh
M., Konno, H., Okazaki, Y., Muramatsu, M. and Hayashizaki, Y.

Normalization and subtraction of cap-trapper-selected cDNAs to
prepare full-length cDNA libraries for rapid discovery of new
genes. Genome Res. 10 (10), 1617-1630 (2000)
wagi, K., Fujiwaka, S., Inoue, K., Togawa, Y., Izawa, M., Ohara, E.,
Watahiki, M., Yoneda, Y., Ishikawa, T., Ozawa, K., Tanaka, T., Matsuura
S., Kawai, J., Okazaki, Y., Muramatsu, M., Inoue, Y., Kira, A. and
Hayashizaki, Y.

RIKEN integrated sequence analysis (RISA) system--384-format
sequencing pipeline with 384 multicapillary sequencer. Genome Res.
10 (11), 1757-1771 (2000)
Konno, H., Fukunishi, Y., Shibata, K., Itoh, M., Carninci, P., Sugahara
Y. and Hayashizaki, Y.
Computer-based methods for the mouse full-length cDNA
encyclopedia: real-time sequence clustering for construction of a
nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)
Please visit our web site (<http://genome.gsc.riken.go.jp>) for
further details.
e mouse tissues.

FEATURES Location/Qualifiers
Source
1..370

/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="F430010H21"
/tissue_type="spleen"
/dev_stage="6 days neonate"
/clone_lib="RIKEN full-length enriched, 6 days neonate
spleen"

BASE COUNT 85 a 116 c 96 g 73 t
ORIGIN

Query Match 13.4%; Score 132.6; DB 10; Length 370;
Best Local Similarity 68.9%; Pred. No. 2e-09;
Matches 202; Conservative 0; Mismatches 79; Indels 12; Gaps 1;

QY 44 GCCCTACACCCAGGAGGATGGGGCTCCCTGGGCTGTCTGTGGCGTGGCTGC 103
DB 90 GCCCAACCTCAGACGATGGCGCTCCTTCAGTCTCTCTTTGCCATGCTGGCTGC 149


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QY 104 CAGCAGCTTCTCAAGCAGCGGAGGAGAAATTACCCCTGTGGTCTCCATTGCTCCACAA 163
Db 150 CTGTGGCTTCTCAGAGAGCAGACAGAGGCATCA-----CCATTGCTCCACAA 197
QY 164 AGTCTCTGGAAGTTTTCCTCCAAAGCGCGCTGGTGCTCATAACTGCTGTGTCACCCAGCC 223
Db 198 AGTACTGGAAGTTTATCCCAAGCGGAGGGTGCTTATTAACCTGCGATGCCCTCAGGC 257
QY 224 ACCACGCCCATCACCTATTCCCTCTGTGTGGAACAAGAACATCAAGGTGGCCCAAGAGGT 283
Db 258 GTCCAGGCCCATCACATACTCTCTCTGCTAGCGAGGTATCTCTGTGGCAAAAAGGT 317
QY 284 GTGGAAGACCCAGCAGCGCGCTCTCTCACTCACTCACTCACTCACTCACTCACTCACTCA 336
Db 318 TGTGATGACTCGTCCGCGGCTCTCTCACTCACTCACTCACTCACTCACTCACTCACTCACT 370

RESULT 34
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LOCUS 260980 MARC 2P1G Sus scrofa cDNA 5', mRNA linear EST 01-DEC-2000
DEFINITION
ACCESSION BF443482
VERSION BF443482.1 GI:11503574
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
AUTHORS Fahrrenkrug,S.C., Smith,T.P.L., Freking,B.A., Cho,J., White,J.,
Vallet,J., Wise,T., Rohrer,G.A., Pertea,G., Sultana,R., Quackenbush
J. and Keefe,J.W.
Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
Mamm. Genome 13 (8), 475-478 (2002)
2213789
MEDLINE 12226715
PUBMED
COMMENT Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -minmatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCCTCCAGTCACGACG
Plate: 93 row: K column: 24
Seq primer: ATTAGTGACACTATAG.
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Location/Qualifiers
1..420
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 2P1G"
/note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, ovary,
endometrium, hypothalamus, pituitary, and placenta."
BASE COUNT 99 a 109 c 130 g 82 t
ORIGIN
Query Match 9.4%; Score 93.2; DB 10; Length 420;
Best Local Similarity 71.1%; Pred. No. 0.00064;
Matches 138; Conservative 0; Mismatches 53; Indels 3; Gaps 1;

QY 428 GTCCAGCCAGTGTCTGAGCTCGGGCCCACTTCACTCTGCAGGACAGAGG---GGCAGG 484
Db 223 GCCAGAGCCCGTGTCCAGCTGTGTGTTAACTTCATCTCTGCTGGAGCGGCGCGCTCGGG 282

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QY 485 CCCACGGGTGGAGATCTGCAGCGCTCTCGGCGAGCCCACTATCAACACAGCCT 544
Db 283 CCCTTGGGTGAAGGTGTCTGTGTGGCATCTCTGGGCGAGCCCACTACCTACAGTCT 342
QY 545 GATCGGGAAGATGGCAGGTCCACCTGCGAGCAGACCATGCCACAGCAGCCTGCCA 604
Db 343 TGTGCGGAGGAGCAGTGGCCACCTATGAAGCAGACACAGAACTACAGGAAGCTGCCA 402
QY 605 CTTCTCCTTCTCTGC 618
Db 403 CTTCTCCTTCTCTGC 416

RESULT 35
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LOCUS 258654 MARC 2P1G Sus scrofa cDNA 5', mRNA linear EST 01-DEC-2000
DEFINITION
ACCESSION BF442092
VERSION BF442092.1 GI:11502184
KEYWORDS EST.
SOURCE Sus scrofa (pig)
ORGANISM
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
Mammalia; Eutheria; Cetartiodactyla; Suina; Suidae; Sus.
REFERENCE
AUTHORS Fahrrenkrug,S.C., Smith,T.P.L., Freking,B.A., Cho,J., White,J.,
Vallet,J., Wise,T., Rohrer,G.A., Pertea,G., Sultana,R., Quackenbush
J. and Keefe,J.W.
Porcine gene discovery by normalized cDNA-library sequencing and
EST cluster assembly
Mamm. Genome 13 (8), 475-478 (2002)
2213789
MEDLINE 12226715
PUBMED
COMMENT Contact: Smith TPL
USDA, ARS, US Meat Animal Research Center
PO Box 166, Clay Center, NE 68933-0166, USA
Tel: 402 762 4366
Fax: 402 762 4390
Email: smith@email.marc.usda.gov
Single pass sequencing. Bases called and alt trimmed with phred
v0.980904.e. Vector identified by cross_match with the -minscore 18
and -minmatch 12 options.
PCR Primers
FORWARD: AGGAACAGCTATGACCAT
BACKWARD: GTTTCCTCCAGTCACGACG
Plate: 88 row: H column: 8
Seq primer: ATTAGTGACACTATAG.
FEATURES
Location/Qualifiers
1..503
/organism="Sus scrofa"
/mol_type="mRNA"
/db_xref="taxon:9823"
/tissue_type="pooled"
/lab_host="DH10B"
/clone_lib="MARC 2P1G"
/note="Vector: pCMV SPORT6; Site 1: NotI; Site 2: SalI;
Library made from pooled tissue from testis, ovary,
endometrium, hypothalamus, pituitary, and placenta."
BASE COUNT 104 a 160 c 160 g 79 t
ORIGIN
Query Match 9.3%; Score 92.6; DB 10; Length 503;
Best Local Similarity 70.8%; Pred. No. 0.00072;
Matches 138; Conservative 0; Mismatches 54; Indels 3; Gaps 1;

QY 433 AGCCAGTGTCTGAGCTCGGCGCACTTCACTCTGCAGGACAGAGG---GGCAGGCCCA 489
Db 309 AGCCCGTGTCCAGCTGTCTGTGTTAACTTCACTCTGCTGGAGCGGCGCGTGGGCCCTT 368
QY 490 GGGTGGAGATGATCTGCCAGGCGCTCTCGGCGAGCCCACTATCAACACAGCCTGATCG 549
Db 369 GGGTGAAGTGTCTGTGCTGGGCATCTCTCGGCGAGCCCACTACCTACCTACAGTCTTGTG 428

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QY 550 GGAAGGATGGCAGAGTCCACCTGCAGCAGAGACCATGCCAGGCGAGCTGCACACTTCT 609
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
429 GGAGGAGCAGTGGCACCCTACATGAGCAGACACAGACTACAGAGAGTGCACACTTCT 488
QY 610 CTTCTCTGCGGAGCC 624
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||

RESULT 36
LOCUS AA031034 208 bp mRNA linear EST 21-AUG-1996
DEFINITION mi47n05.r1 Soares mouse embryo NbME13.5 14.5 Mus musculus cDNA
clone IMAGE:466713 5', mRNA sequence.
ACCESSION AA031034
VERSION AA031034.1 GI:1501031
KEYWORDS EST.
SOURCE Mus musculus (house mouse)
ORGANISM Mus musculus
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Rodentia; Sciurognathi; Muridae; Murinae; Mus.
1 (bases 1 to 208)
Geisel,S., Kucab,T., Allen,M., Bowles,M., Dietrich,N., Dubuque,T.,
Schellenberg,K., Steptoe,M., Tan,F., Underwood,K., Moore,B.,
Theising,B., Wylie,T., Lennon,G., Soares,B., Wilson,R. and
Waterston,R.
TITLE The WashU-HMI Mouse EST Project
JOURNAL Unpublished
COMMENT Contact: Marra M/Mouse EST Project
WashU-HMI Mouse EST Project
Washington University School
4444 Forest Park Parkway, Box 8501, St. Louis, MO 63108
Tel: 314 286 1800
Fax: 314 286 1810
Email: mouseest@wustl.edu
This clone is available royalty-free through LNL; contact the
IMAGE Consortium (info@image.llnl.gov) for further information.
MGI:280529
Seq primer: -28M13 rev2 from Amerham
High quality sequence stop: 193.
FEATURES
source
location/Qualifiers
1..208
/organism="Mus musculus"
/mol_type="mRNA"
/strain="C57BL/6J"
/db_xref="taxon:10090"
/clone="IMAGE:466713"
/tissue_type="embryo"
/dev_stages="13.5-14.5dpc total fetus"
/lab_host="DH10B"
/clone_lib="Soares mouse embryo NbME13.5 14.5"
/note="Vector: pYT3D-Pac (Pharmacia) with a modified
polylinker; Site_1: Not I; Site_2: Eco RI; 1st strand cDNA
was primed with a Not I - oligo(dT) primer 15',
TGTTACCAATCAGTGGAGCGCGCCGCGAATTTTTTTTTTTTTTTT
T 3', on equal amounts of mRNA from 2 13.5dpc and 2
14.5dpc embryos [total RNA provided by Minoru Ko, Wayne
State Univ., from 2 ]; double-stranded cDNA was ligated to
Eco RI adaptors (Pharmacia), digested with Not I and
cloned into the Not I and Eco RI sites of the modified
pYT73 vector. Library went through one round of
normalization, and was constructed by Bento Soares and
M.Fatima Bonaldo."
BASE COUNT 47 a 61 c 51 g 49 t
ORIGIN
Query Match 8.0%; Score 79.8; DB 9; Length 208;
Best Local Similarity 65.3%; Pred. No. 0.067;
Matches 139; Conservative 0; Mismatches 62; Indels 12; Gaps 1;

QY 77 GCTGTCTCTGTCGGCTGCTGCAGCAGACTTCTCCAGGCGGAGGAGAAAT 136

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Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
7 GCTGTCTCTTTCATGCTGGCTGCCTGTGCGTTCTCAGAGGAGCAGACAGAGGCA- 65
QY 137 TACCCCTGTGCTCCATTGCTTACAAAGTCTGTGAAGTTTTCCTCCAAAGCGCGTGGGT 196
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
66 -----TCACCAATTGCTTACAAAGTACTGGAAGTTTATCCCAAGACCGGAGGTA 114
QY 197 GCTCATATACCTGTGTGCACCCAGCCACCGCCATCACCATTATTCCTCTGTGGAC 256
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
115 GCTTATAACCTGCGATGCCCTGTAGGCGTCCAGCCCATCATACTCTCTCTGGCTAG 174
QY 257 CAAGAACAATCAAGTGGCCAAAGAGTGGTGAA 289
Db ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| ||||| |||||
175 CCGAGGTATCTGTGGTGGCAAAAAGTTGTGCA 207

RESULT 37
LOCUS AI187208/483 bp mRNA linear EST 10-NOV-1998
DEFINITION qf28b11.x1 Soares testis NHT Homo sapiens cDNA clone IMAGE:1751325
3', similar to contains MER27.tl MER32 repetitive element ;, mRNA
sequence.
ACCESSION AI187208
VERSION AI187208.1 GI:3737846
KEYWORDS EST.
SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
REFERENCE Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;
AUTHORS Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.
1 (bases 1 to 483)
TITLE NCI-CGAP http://www.ncbi.nlm.nih.gov/ncicgap.
JOURNAL National Cancer Institute, Cancer Genome Anatomy Project (CGAP),
COMMENT Unpublished
Contact: Robert Strausberg, Ph.D.
Email: cgapbs-remail.nih.gov
cDNA Library Preparation: M. Bento Soares, Ph.D., M. Fatima Bonaldo
, Ph.D.
cDNA Library Arrayed by: Greg Lennon, Ph.D.
DNA Sequencing by: Washington University Genome Sequencing Center
Clone distribution: NCI-CGAP clone distribution information can be
found through the I.M.A.G.E. Consortium/LLNL at:
www-bio.llnl.gov/bbrp/image/image.html
Insert Length: 678 Std Error: 0.00
Seq primer: -40UP from Gibco
High quality sequence stop: 381.
FEATURES
source
location/Qualifiers
1..483
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="IMAGE:1751325"
/sex="male"
/lab_host="DH10B"
/clone_lib="Soares testis NHT"
/note="Vector: pYT3D-Pac (Pharmacia) with a modified
polylinker; Site 1: Not I; Site 2: Eco RI; 1st strand cDNA
was prepared from mRNA obtained from Clontech Laboratories
Inc. and primed with a Not I - oligo(dT) primer [5',
TGTTACCAATCAGTGGAGCGCGCCCAATTTTTTTTTTTTTTTT 3'].
Double-stranded cDNA was ligated to Eco RI adaptors
(Pharmacia), digested with Not I and cloned into the Not I
and Eco RI sites of the modified pYT73 vector. Library
went through one round of normalization to Cots5, and was
constructed by Bento Soares and M. Fatima Bonaldo."
BASE COUNT 133 a 103 c 78 g 169 t
ORIGIN
Query Match 5.8%; Score 57.8; DB 9; Length 483;
Best Local Similarity 67.9%; Pred. No. 56;
Matches 112; Conservative 0; Mismatches 47; Indels 6; Gaps 2;

QY 829 AGTTCAGGAGCGCAAGAGCAGCCCATGTAGTAATGAACCGTCCAGAGGC--CAAGCAC 886

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Db      174 GGGAGATAGTGGAGGATACAAAGTAGCAGATATAGATGATGAGCAAGTC---TAGA 118
QY      880 CAAGCAGCGCAGAGACTGCGAGGCCATCAGCGTGTCACCTGTTTCGTAATTTGGAGTTCATGCA 939
Db      117 GAAATTAATGTACATGAGGACCTCTAGGTAAACACAATTTGATGTATGTTGGAATTCATGCT 58
QY      940 AAATGAGTGTGTTTACGTCTCTTGGCCACAAAAAAGAAAAA 982
Db      57 AAAGGAGTACAATTTAGTCTCTTGGCAACAAAAACAAGAAA 15

RESULT 40
LOCUS   BI063151
DEFINITION IL3-UT0117-300301-512-B06 UT0117 Homo sapiens cDNA, mRNA sequence. EST 15-JUN-2001
ACCESSION BI063151
VERSION    BI063151.1 GI:14470678
KEYWORDS  EST.
SOURCE    Homo sapiens (human)
ORGANISM  Homo sapiens
REFERENCE 1 (bases 1 to 606)
AUTHORS   Dias Neto,E., Garcia Correa,R., Verjovski-Almeida,S., Briones,M.R., Nagai,M.A., da Silva,W. Jr., Zago,M.A., Bordin,S., Costa,F.F., Goldman,G.H., Carvalho,A.F., Matsukuma,A., Baia,G.S., Simpson,D.H., Brunstein,A., deoliveira,P.S., Bucher,P., Jongeneel,C.V., O'Hare,M.J., Soares,F., Brentani,R.R., Reis,L.F., de Souza,S.J. and Simpson,A.J.
TITLE     Shotgun sequencing of the human transcriptome with ORF expressed sequence tags
JOURNAL   Proc. Natl. Acad. Sci. U.S.A. 97 (7), 3491-3496 (2000)
MEDLINE   20202663
PUBMED    10737800
COMMENT   Contact: Simpson A.J.G.
          Laboratory of Cancer Genetics
          Ludwig Institute for Cancer Research
          Rua Prof. Antonio Prudente 109, 4 andar, 01509-010, Sao Paulo-SP, Brazil
          Tel: +55-11-2704922
          Fax: +55-11-2707001
          Email: asimpson@ludwig.org.br
          This sequence was derived from the FAPESP/LICR Human Cancer Genome project. This entry can be seen in the following URL
          (http://www.ludwig.org.br/scripts/gethtml2.pl?l1=IL3&t2=IL3-UT0117-300301-512-B06&t3=2001-03-30&t4=1)
          Seq primer: puc 18 forward
          High quality sequence stop: 502.
          Location/Qualifiers
            1..606
              /organism="Homo sapiens"
              /mol_type="mRNA"
              /db_xref="taxon:9606"
              /dev_stage="Adult"
              /clone_lib="UT0117"
              /note="Organ: uterus_tumor; Vector: puc18; Site_1: SmaI; Site_2: SmaI; A mini-library was made by cloning products derived from ORESTES PCR (U.S. Letters Patent application No. 196,716 - Ludwig Institute for Cancer Research) profiles into the pUC 18 vector. Reverse transcription of tissue mRNA and cDNA amplification were performed under low stringency conditions."
            124 a 167 c 143 g 172 t

BASE COUNT 124 a 167 c 143 g 172 t
ORIGIN

Query Match
Best Local Similarity 5.2%; Score 51.2; DB 12; Length 606;
Matches 113; Conservative 0; Mismatches 63; Indels 4; Gaps 2;

QY      813 AGGATAGGGAATGGGAGGTGAGGAGCGAAGCAGAGCGGATGTAGAAATGAAACCGGCC 872
Db      540 AGGAAACGGGAGATGTAGTGAAGAGCGCAGAGAGGCGAGATATGCAAGATGCATCGCC 481

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QY      873 AGAGAGCAAGCAGCGGAGGAGTCTGAGGCGCATCAGCTGTCACCTGTCGTTGTTTGGAGT 932
Db      480 AGAG-GCCTAATGTGCATGAGGACTCTCTAGTCAACACATTTGA---ATTATACCGGGAT 425
QY      933 TCATGCAAAATGAGTGTGTTTACGTCTCTTGGCCAGAAAAAAGAAAAAAGAAAAA 992
Db      424 TTCTGTAGAGACAGTAGATTTTAGGAGCTCTTGCATATTGAAAAAAGAAAAAGGTAAA 365

RESULT 41
LOCUS   BY560588
DEFINITION BY560588 RIKEN full-length enriched, activated spleen Mus musculus cDNA clone F830028K04 3', mRNA sequence.
ACCESSION BY560588
VERSION    BY560588.1 GI:26895770
KEYWORDS  EST.
SOURCE    Mus musculus (house mouse)
ORGANISM  Mus musculus
REFERENCE 1 (bases 1 to 443)
AUTHORS   Okazaki,Y., Furuno,M., Kasukawa,T., Adachi,J., Bono,H., Kondo,S., Nikaido,I., Osato,N., Saito,R., Suzuki,H., Yamanaka,I., Kiyosawa,H., Yagi,K., Tomaru,Y., Hasegawa,Y., Nogami,A., Schonbach,C., Gojobori,T., Baldarelli,R., Hill,D.P., Bult,C., Hume,D.A., Quackenbush,J., Schriml,L.M., Kanapin,A., Matsuoka,H., Batalov,S., Beisel,K.W., Blake,J.A., Bradt,D., Brusci,V., Chothia,C., Corbani,L.E., Cousins,S., Dalla,E., Dragani,T.A., Fletcher,C.F., Forrest,A., Frazer,K.S., Gaasterland,T., Gariboldi,M., Gissi,C., Godzik,A., Gough,J., Grammond,S., Gustinchin,S., Hirokawa,N., Jackson,I.J., Jarvis,E.D., Kanai,A., Kawaji,H., Kawasawa,Y., Kedzierski,R.M., King,B.L., Konagaya,A., Kurochkin,I.V., Lee,Y., Lenhard,B., Lyons,P.A., Maglott,D.R., Maltais,L., Marchionni,L., McKenzie,L., Miki,H., Nagashima,T., Numata,K., Okido,T., Pavan,W.J., Pertea,G., Pesole,G., Petrovsky,N., Pillai,R., Pontius,J.U., Qi,D., Ramachandran,S., Ravasi,T., Reed,J.C., Reed,D.J., Reid,J., Ring,B.Z., Ringwald,M., Sandelin,A., Schneider,C., Semple,C.A., Setou,M., Shimada,K., Sultana,R., Takenaka,Y., Taylor,M.S., Teasdale,R.D., Tomita,M., Verardo,R., Wagner,L., Wahlestedt,C., Wang,Y., Watanabe,Y., Wells,C., Wilming,L.G., Wyshaw-Boris,A., Yanagisawa,M., Yang,I., Yang,L., Yuan,Z., Zavolan,M., Zhu,Y., Zimmer,A., Carninci,P., Hayatsu,N., Hirozane-Kishikawa,T., Konno,H., Nakamura,M., Sakazume,N., Sato,K., Shiraki,T., Waki,K., Kawai,J., Aizawa,K., Arakawa,T., Fukuda,S., Hara,A., Hashizume,W., Imorani,K., Ishii,Y., Itoh,M., Kagawa,I., Miyazaki,A., Sakai,K., Sasaki,K., Shibata,K., Shinagawa,A., Yasunishi,A., Yoshino,M., Waterston,R., Lander,E.S., Rogers,J., Birney,E. and Hayashizaki,Y.
TITLE     Analysis of the mouse transcriptome based on functional annotation of 60,770 full-length cDNAs
JOURNAL   Nature 420, 563-573 (2002)
MEDLINE   22354683
PUBMED    12466851
COMMENT   Contact: Yoshihide Hayashizaki
          Laboratory for Genome Exploration Research Group, RIKEN Genomic Sciences Center (GSC), Yokohama Institute
          The Institute of Physical and Chemical Research (RIKEN)
          1-7-22 Suehiro-cho, Tsurumi-Ku, Yokohama, Kanagawa 230-0045, Japan
          Tel: 81-45-503-9222
          Fax: 81-45-503-9216
          Email: genome-res@gsc.riken.go.jp,
          URL: http://genome.gsc.riken.go.jp/
          Aizawa,K., Akimura,T., Arakawa,T., Itoh,M., Kawai,J., Konno,H., Miyazaki,A., T., Imotani,K., Ishii,Y., Itoh,M., Nomura,K., Numazaki,R., Ohno,M., Sakai,K., Murata,M., Nakamura,M., Nomura,K., Shibata,K., Shiraki,T., Tagami,Sakazume,N., Sasaki,D., Sato,K., Shibata,K., Shiraki,T., Tagami,M., Waki,K., Watahiki,A., Muramatsu,M. and Hayashizaki,Y. Direct Submission
          Computational Analysis of Full-Length Mouse cDNAs Compared with Human Genome Sequences Mamm. Genome. 12, 673-677 (2001)
          Normalization and subtraction of cap-trapper-selected cDNAs to prepare full-length cDNA libraries for rapid discovery of new genes. Genome Res. 10 (10), 1617-1630 (2000)

```

RIKEN integrated sequence analysis (RISA) system--384-format sequencing pipeline with 384 multipillar sequencer. Genome Res. 10 (11), 1757-1771 (2000)

Computer-based methods for the mouse full-length cDNA encyclopedia: real-time sequence clustering for construction of a nonredundant cDNA library. Genome Res. 11 (2), 281-289 (2001)

cDNA library was prepared and sequenced in Mouse Genome Encyclopedia Project of Genome Exploration Research Group in Riken Genomic Sciences Center and Genome Science Laboratory in RIKEN. Division of Experimental Animal Research in Riken contributed to prepare mouse tissues.

Tissues were provided by Dr. John Todd (Dept. of Medical Genetics Wellcome Trust Centre for Molecular Mechanisms in Disease Wellcome Trust/MRC building Addenbrookes Hospital Cambridge) whose assistance we gratefully acknowledge.

Please visit our web site (<http://genome.gsc.riken.go.jp>) for further details.

FEATURES

source

Location/Qualifiers

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1. .443
/organism="Mus musculus"
/mol_type="mRNA"
/strain="NOD"
/db_xref="taxon:10090"
/clone="F830028K04"
/tissue_type="activated spleen"
/clone_lib="RIKEN full-length enriched, activated spleen"
BASE COUNT 97 a 128 c 117 g 101 t
ORIGIN
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Query Match 5.1%; Score 50.8; DB 13; Length 443;
Best Local Similarity 62.7%; Pred. No. 5.6e+02;
Matches 79; Conservative 0; Mismatches 47; Indels 0; Gaps 0;

QY 597 CTTGCCAATCTCTCTCCGCGAGCCAGACATCGAGTGTCTGTCGCGAGGTGCA 656
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
1 CAGACCAATCTCTCTCCGCGAGCCAGACATCGAGTGTCTGTCGCGAGGTGCA 60

QY 657 AACACGCCAATCTCCAGCAGCGCCCTCACAGTGTGCCCCCGAGTGACAGAG 716
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
61 AACGATGTGGTGTGACAGCAGTGTGCGCGATCCCGTCCCGAGGCGCGAGCCAAG 120

QY 717 ATGGAG 722
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
121 CTGGTG 126
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RESULT 42

CNS0091P

LOCUS

DEFINITION

Drosophila melanogaster genome survey sequence TET3 end of BAC # BACR19D16 of RPI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Drosophila melanogaster (fruit fly)
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 925)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr
Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osoegawa and Aaron Mammosier in Pieter de Jong's laboratory in the Department of

Cancer Genetics at the Roswell Park Cancer Institute in Buffalo, NY. The library is named RPI-98 and was constructed by partial EcoRI digestion of Drosophila DNA provided by the BDGP from the isogenic strain Y2; cn bw sp, the same strain used for the BDGP's pl and EST libraries. A more detailed description of the library and how to order individual BAC clones, the entire library, or filters for hybridization from the BACPAC Resource Center can be found at http://bacpac.med.buffalo.edu/drosophila_bac.htm.

FEATURES

source

Location/Qualifiers

1. .925

/organism="Drosophila melanogaster"

/mol_type="genomic DNA"

/db_xref="taxon:7227"

/clone="BACR19D16"

/clone_lib="RPI-98"

/note="end : TET3"

BASE COUNT 120 a 61 c 61 g 172 t 511 others

ORIGIN

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Query Match 4.9%; Score 49; DB 29; Length 925;
Best Local Similarity 15.1%; Pred. No. 7.2e+02;
Matches 47; Conservative 138; Mismatches 127; Indels 0; Gaps 0;
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QY 292 CCACGAGCGCGCTCTCAACCTCAAGTCACTCACTCACTCACTCACTCACTCA 351
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
572 CCSCSCSCSSSCSCCHCCSCSCSSYCCSSSBSSKCSSTSBSCSSKSVCGTSC 631

QY 352 CTTACTTCTGCGGGCGCTCTCCACTCAGGTGCCATGTGCACAGTCCAGCTACAGA 411
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
632 SSSSSSCSSSSSTSSSTSSSTSSKSSSSSSSSSSSYTTSKTSASGGSWSAGGSSS 691

QY 412 TGCATGTGGAGTGTGTGTCACAGCAGTGTCTGAGTGTGCGGCCCACTTCTCTGACG 471
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
692 TGTSTSSSSSTSTSSSVSGSKSTBSGSSBSSSSSSSTSSBBSCTSTSSSSSS 751

QY 472 ACAGAGGGCAGCGCCCGAGGTGGAGATCATGTGCCAGCGCTCTCGGCGACGCCACTA 531
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
752 STSSSTCTCTCCCTCCSSSYSSSTSSSTSSSTSSSTSSSTSSSTSSSTSSSTSSSTSS 811

QY 532 TCACCAACAGCTGTATCGGAGAGATGGCAGTCCACCTCGCAGCAGACCATGCGACA 591
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
812 CTCCTSTYMBCTSTSTCGSSSSSSSGKGVTKCGCGGSSSTNGBTSSACSSSSSSCS 871

QY 592 GGCAGCTGCCA 603
Db ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| ||| |||
872 SSSVSSSKSSA 883
```

RESULT 43

CNS0052P

LOCUS

DEFINITION

Drosophila melanogaster genome survey sequence TET3 end of BAC # BACR11P16 of RPI-98 library from Drosophila melanogaster (fruit fly), genomic survey sequence.

ACCESSION

VERSION

KEYWORDS

SOURCE

ORGANISM

Drosophila melanogaster (fruit fly)
Eukaryota; Metazoa; Arthropoda; Hexapoda; Insecta; Pterygota; Neoptera; Endopterygota; Diptera; Brachycera; Muscomorpha; Ephydroidea; Drosophilidae; Drosophila.
1 (bases 1 to 844)

REFERENCE

AUTHORS

TITLE

JOURNAL

COMMENT

Submitted (02-JUN-1999) Genoscope - Centre National de Sequencage : BP 191 91006 EVRY cedex - FRANCE (E-mail : segref@genoscope.cns.fr)
- Web : www.genoscope.cns.fr
Determination of this BAC-end sequence was carried out as part of a collaboration with the Berkeley Drosophila Genome Project (BDGP). The BDGP is constructing a physical map of the Drosophila melanogaster genome using these BACs. For further information please see <http://www.fruitfly.org> The BDGP Drosophila melanogaster BAC library was prepared by Kazutoyo Osoegawa and Aaron Mammosier in Pieter de Jong's laboratory in the Department of

COMMENT

Contact: Genoscope
Genoscope - Centre National de Sequençage
BP 191 91006 EVRY cedex - France
Email: segref@genoscope.cns.fr, Web : www.genoscope.cns.fr
Library was constructed by Life Technologies, a division of
Invitrogen. Contact : Feng Liang Email : liang@lifetech.com URL :
<http://fulllength.invitrogen.com/> Invitrogen Corporation 1600
Faraday Avenue Genoscope sequence ID : CS0D1070AG02NP1.

FEATURES

```

library location: Genoscope sequence ID : CS0D1070AG02NP1.
Accession/Qualifiers
1..974
/organism="Homo sapiens"
/mol_type="mRNA"
/db_xref="taxon:9606"
/clone="CS0D1070YM03"
/tissue_type="PLACENTA COT 25-NORMALIZED"
/clone_lib="Homo sapiens PLACENTA COT 25-NORMALIZED"
/notes="1st strand cDNA was primed with a NotI-oligo(dT)
primer.. Five prime end enriched, double-strand cDNA was
digested with Not I and cloned into the Not I and EcoR V
sites of the pCMVSPORT 6 vector. Library was normalized."
69 a 235 c 80 q 310 t 280 others

```

[illegible]

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Query Match      4.8%; Score 47.2; DB 13; Length 974;
Best Local Similarity 25.8%; Pred. No. 1.3e+03;
Matches 85; Conservative 95; Mismatches 150;
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Qy	813	AGGATGGGGAATGGGGAGGTCAGAGGACGCAAAAGCAGCAGCCATGTAGAAATGAACCGTC	872
Db	393	AGGAATGGGGAGATGATGTCAGGGGTCAAAATAGCAGATATGTAGAAATAACAGGTC	334

[illegible]

QY	873	AGAGAGCCAAAGCACGGCAGAGACTGCAGGGCCATCAGCOTGCATGTTTCGTATTGGAGT	932
Db	333	TAGAAATGTAATGTATTAACATCG-GGCAGGATTAACATAAATTCGCTGTATTTGGCAT	275

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DEFINITION	clone CS0D1015YB03 3-PRIME, mRNA sequence.		
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DEFINITION	BX356664 Homo sapiens PLACENTA COT 25-NORMALIZED Homo sapiens CDNA		

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SOURCE Homo sapiens (human)

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QY 963 TTGCCACAAAAAIAAAAAAAAAAAAAAAAAA 992

ORGANISM

Db 609 ANAAAAAAAAAAAAAAAAAAAAA 580

REFERENCE	AUTHORS	TITLE	JOURNAL
1 (bases 1 to 1201)	Li, W.B., Gruber, C., Jessee, J. and Polayes, D.	Full-length cDNA libraries and normalization	Unpublished

RESULTS	LOCUS	748 bp	DNA	linear	GSS 13-MAR-2003
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DEFINITION		mcv51f02.g1	HFOSMID007	Homo sapiens genomic, genomics survey	

CONTACT : Genoscope
Genoscope - Centre National de Sequencage
BP 191 91006 EVRY cedex - France
Email : secref@genoscope.cnr.fr Web : www.genoscope.cnr.fr

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VERSION BZ772737.1 GI:28946421
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Library was constructed by Life Technologies, a division of Invitrogen. Contact : Feng Liang Email : liang@lifetech.com URL : <http://fulllength.invitrogen.com/> Invitrogen Corporation 1600

SOURCE Homo sapiens (human)
ORGANISM Homo sapiens
Eukaryota; Metazoa; Chordata; Craniata; Vertebrata; Euteleostomi;

FEATURES
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Location/Qualifiers
organization="Homo sapiens"
Faraday Avenue Genoscope sequence ID: CS0D1015CA02N2P1.

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AUTHORS

Cook, L., Deteaunty, K., Fewell, G., Fulton, L., Magrini, V., Mardis, E.
1 (bases 1 to 748)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

Cook, L., Deteaunty, K., Fewell, G., Fulton, L., Magrini, V., Mardis, E.
1 (bases 1 to 748)
Mammalia; Eutheria; Primates; Catarrhini; Hominidae; Homo.

Miner, T., Nash, W., Williams, D. and Wilson, R.K.

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TITLE	JOURNAL	COMMENT
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primer. Five prime end enriched, double-strand cDNA was

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Genetic Sequencing Center
Washington University School of Medicine
Email: submissions@watson.wustl.edu

BASE COUNT 116 a 88 c 93 g 398 t 506 others

[illegible]

[illegible]

GenCore version 5.1.6
Copyright (c) 1993 - 2004 Compugen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 9, 2004, 13:04:22 | Search time 164 Seconds
(without alignments)
2669.830 Million cell updates/sec

Title: US-09-990-726-222

Perfect score: 992

Sequence:

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Scoring table: IDENTITY_NUC

Gapop 10.0, Gapext 1.0

Searched: 569978 seqs, 220691566 residues

Total number of hits satisfying chosen parameters: 1139956

Minimum DB seq length: 0

Maximum DB seq length: 2000000000

Post-processing: Minimum Match 0%

Maximum Match 100%

Listing first 150 summaries

Database :

Issued Patents NA:*

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

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ALIGNMENTS

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; APPLICANT: Baker, Kevin P.
; APPLICANT: Botstein, David
; APPLICANT: Desnovers, Luc
; APPLICANT: Eaton, Dan L.
; APPLICANT: Ferrara, Napoleone
; APPLICANT: Fong, Sherman
; APPLICANT: Gerber, Hanspeter
; APPLICANT: Gerritsen, Mary E.
; APPLICANT: Goddard, Audrey
; APPLICANT: Godowski, Paul J.
; APPLICANT: Grimaldi, J. Christopher
; APPLICANT: Gurney, Austin L.

; APPLICANT: Kljavin, Ivar J.
; APPLICANT: Napier, Mary A.
; APPLICANT: Pan, James
; APPLICANT: Paoni, Nicholas F.
; APPLICANT: Roy, Margaret Ann
; APPLICANT: Stewart, Timothy A.
; APPLICANT: Tumas, Daniel
; APPLICANT: Watanabe, Colin K.
; APPLICANT: Williams, P. Mickey
; APPLICANT: Wood, William I.
; APPLICANT: Zhang, Zemin
; TITLE OF INVENTION: Secreted and Transmembrane Polypeptides and Nucleic
; FILE OF INVENTION: Acids Encoding the Same
; FILE REFERENCE: P2730P1C13
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;; PRIOR FILING DATE: 1998-06-24

;; PRIOR APPLICATION NUMBER: 60/090542
;; PRIOR FILING DATE: 1998-06-24
;; PRIOR APPLICATION NUMBER: 60/090557
;; PRIOR FILING DATE: 1998-06-24
;; PRIOR APPLICATION NUMBER: 60/090676
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090678
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090690
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090694
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090695
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090696
;; PRIOR FILING DATE: 1998-06-25
;; PRIOR APPLICATION NUMBER: 60/090862
;; PRIOR FILING DATE: 1998-06-26
;; PRIOR APPLICATION NUMBER: 60/090863
;; PRIOR FILING DATE: 1998-06-26
;; PRIOR APPLICATION NUMBER: 60/091360
;; PRIOR FILING DATE: 1998-07-01
;; PRIOR APPLICATION NUMBER: 60/091478
;; PRIOR FILING DATE: 1998-07-02
;; PRIOR APPLICATION NUMBER: 60/091544
;; PRIOR FILING DATE: 1998-07-01
;; PRIOR APPLICATION NUMBER: 60/091519
;; PRIOR FILING DATE: 1998-07-02
;; PRIOR APPLICATION NUMBER: 60/091626
;; PRIOR FILING DATE: 1998-07-02
;; PRIOR APPLICATION NUMBER: 60/091633
;; PRIOR FILING DATE: 1998-07-02
;; PRIOR APPLICATION NUMBER: 60/091978
;; PRIOR FILING DATE: 1998-07-07
;; PRIOR APPLICATION NUMBER: 60/091982
;; PRIOR FILING DATE: 1998-07-07
;; PRIOR APPLICATION NUMBER: 60/092182
;; PRIOR FILING DATE: 1998-07-09

Query Match

100.0%; Score 992; DB 4; Length 992;

Best Local Similarity 100.0%; Pred. No. 1.4e-243;

Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCACGAGCCAGAACTAGAGGTTCTCACTCCCGAGCAGAGGCCCTACACCCACCGAG 60
Db 1 GGCACGAGCCAGAACTAGAGGTTCTCACTCCCGAGCAGAGGCCCTACACCCACCGAG 60
QY 61 GCATGGGGCTCCCTGGGCTGTTCTGCTTGGCGGTGCTGGCTCCAGCAGCTTCTCCAGG 120
Db 61 GCATGGGGCTCCCTGGGCTGTTCTGCTTGGCGGTGCTGGCTCCAGCAGCTTCTCCAGG 120
QY 121 CACGGGAGGAGAAATTACCCCTGCTGCTTCCATTGCTCCTACAAAGTCTGGAAGTTTCC 180
Db 121 CACGGGAGGAGAAATTACCCCTGCTGCTTCCATTGCTCCTACAAAGTCTGGAAGTTTCC 180
QY 181 CCAAAGGCGGTGGGTGCTCATAACTGTGTGACCCCGAGCCACCCGCCCCATCACT 240
Db 181 CCAAAGGCGGTGGGTGCTCATAACTGTGTGACCCCGAGCCACCCGCCCCATCACT 240
QY 241 ATTCCCTCTGTGGAAACCAAGAACATCAAGGTGGCCAAAGGTGGTGAAGACCCACGAGC 300
Db 241 ATTCCCTCTGTGGAAACCAAGAACATCAAGGTGGCCAAAGGTGGTGAAGACCCACGAGC 300
QY 301 CGGCGCTCTTCAACTCAACGTCACTCAAGTCCAGTCCAGCTGCTCACTACTTCT 360
Db 301 CGGCGCTCTTCAACTCAACGTCACTCAAGTCCAGTCCAGCTGCTCACTACTTCT 360
QY 361 GCGGGCGTCTCCACCTCAGTGCCCATGTGGAGCAGTCCAGGCTACAGATGCACTGGG 420
Db 361 GCGGGCGTCTCCACCTCAGTGCCCATGTGGAGCAGTCCAGGCTACAGATGCACTGGG 420
QY 421 AGCTGTGTCCAAGCAGTGTCTGAGTGGCGGCAATTCATCTGTGAGCAGAGGGG 480
Db 421 AGCTGTGTCCAAGCAGTGTCTGAGTGGCGGCAATTCATCTGTGAGCAGAGGGG 480

Db 421 AGCTGTGTCAGCCAGTGTCTGAGCTGCGGGCCAACTTCACTCTGCAGGACAGAGGG 480
QY 481 CAGGCCCCAGGGTGGAGATGATCTGCCAGGCTCTCGGGCAGCCACCTATCAACA 540
Db 481 CAGGCCCCAGGGTGGAGATGATCTGCCAGGCTCTCGGGCAGCCACCTATCAACA 540
QY 541 GCCTGATCGGGAAGATGGGAGTGGTCCCTCAGCAGAGACCATGCCACAGGCGCTG 600
Db 541 GCCTGATCGGGAAGATGGGAGTGGTCCCTCAGCAGAGACCATGCCACAGGCGCTG 600
QY 601 CCAACTTCTCTCTCTCGGAGCCAGACATCGACTGCTTCTGGTCCAGGCTCAACA 660
Db 601 CCAACTTCTCTCTCTCGGAGCCAGACATCGACTGCTTCTGGTCCAGGCTCAACA 660
QY 661 AGCCCAATGTCAGCAGAGCCCTCACAGTGGTGGCCCCAGGTGGTGCACAGAGATGG 720
Db 661 AGCCCAATGTCAGCAGAGCCCTCACAGTGGTGGCCCCAGGTGGTGCACAGAGATGG 720
QY 721 AGGACTGGCAGGTCCTCTGGAGGCCCCCATCTTGGCTTGGCGCTCTACAGGACCCC 780
Db 721 AGGACTGGCAGGTCCTCTGGAGGCCCCCATCTTGGCTTGGCGCTCTACAGGACCCC 780
QY 781 GCGCTCTGAGTCAAGCAGAGTCTGGGGGTTTCAGGATAGGGAATGGGGAGTTCAGGAC 840
Db 781 GCGCTCTGAGTCAAGCAGAGTCTGGGGGTTTCAGGATAGGGAATGGGGAGTTCAGGAC 840
QY 841 GCAAGCAGAGCCATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 900
Db 841 GCAAGCAGAGCCATGATGATGATGATGATGATGATGATGATGATGATGATGATGATG 900
QY 901 GGCATCAGCGTGCCTCTGATTTGGAGTTCATGCAAAATGAGTGTGTTAGTCTGC 960
Db 901 GGCATCAGCGTGCCTCTGATTTGGAGTTCATGCAAAATGAGTGTGTTAGTCTGC 960
QY 961 TCTTGGCCACAAAAA992
Db 961 TCTTGGCCACAAAAA992

RESULT 2

US-08-781-891-79/c
; Sequence 79, Application US/08781891
; Patent No. 6090620
; GENERAL INFORMATION:
; APPLICANT: Fu, Ying-Hui
; APPLICANT: Yu, Chang-En
; APPLICANT: Oshima, Junko
; APPLICANT: Mulligan, John T.
; APPLICANT: Schellenberg, Gerald D.
; TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
; TITLE OF INVENTION: WERNER'S SYNDROME
; NUMBER OF SEQUENCES: 209
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: SEED AND BERRY LLP
; STREET: 6300 Columbia Center, 701 Fifth Avenue
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/781,891
; FILING DATE: 27-DEC-1996
; CLASSIFICATION: 800
; ATTORNEY/AGENT INFORMATION:
; NAME: No. 6090620tenburg Ph.D., Carol
; REGISTRATION NUMBER: 39,317
; REFERENCE/DOCKET NUMBER: 240052.419
; TELECOMMUNICATION INFORMATION:

; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 79:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 87350 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; US-08-781-891-79

Query Match 4.7%; Score 47; DB 3; Length 87350;
Best Local Similarity 78.9%; Pred. No. 0.045;
Matches 56; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 922 GTATTGGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTTCGCCACAAAAA981
Db 18994 GTATTGGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTTCGCCACAAAAA981
QY 982 AAAAAAAAAA992
Db 18834 ATGTTTAAACAA 18824

RESULT 3

US-09-618-166-79/c
; Sequence 79, Application US/09618166
; Patent No. 6583112
; GENERAL INFORMATION:
; APPLICANT: Fu, Ying-Hui
; APPLICANT: Yu, Chang-En
; APPLICANT: Oshima, Junko
; APPLICANT: Mulligan, John T.
; APPLICANT: Schellenberg, Gerald D.
; TITLE OF INVENTION: GENE AND GENE PRODUCTS RELATED TO
; TITLE OF INVENTION: WERNER'S SYNDROME
; NUMBER OF SEQUENCES: 209
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Seed Intellectual Property Law Group
; STREET: 701 Fifth Avenue, Suite 6300
; CITY: Seattle
; STATE: Washington
; COUNTRY: USA
; ZIP: 98104-7092
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent in Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/618,166
; FILING DATE: 17-Jul-2000
; CLASSIFICATION: <Unknown>
; ATTORNEY/AGENT INFORMATION:
; NAME: Mcmasters, David D.
; REGISTRATION NUMBER: 33,963
; REFERENCE/DOCKET NUMBER: 240052.419C1
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (206) 622-4900
; TELEFAX: (206) 682-6031
; INFORMATION FOR SEQ ID NO: 79:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 87350 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; SEQUENCE DESCRIPTION: SEQ ID NO: 79:
; US-09-618-166-79

Query Match 4.7%; Score 47; DB 4; Length 87350;
Best Local Similarity 78.9%; Pred. No. 0.045;
Matches 56; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 922 GTATTGGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTTCGCCACAAAAA981

Db 18894 GTATTGGGATTCCTGCTAAATGAGTGGATTTCAGCTGCTTTGCCACAAAAACAAAA 18835
QY 982 AAAAAAAAAA 992
Db 18834 ATGGTTAACAA 18824

RESULT 4

US-09-791-211-3/C
Sequence 3, Application US/09791211
Patent No. 6448080
GENERAL INFORMATION:
APPLICANT: Donna T. Ward
APPLICANT: Andrew T. Watt
TITLE OF INVENTION: ANTISENSE MODULATION OF WEN EXPRESSION
FILE REFERENCE: RTS-0205
CURRENT APPLICATION NUMBER: US/09/791,211
CURRENT FILING DATE: 2001-02-23
NUMBER OF SEQ ID NOS: 90
SEQ ID NO 3
LENGTH: 87543
TYPE: DNA
ORGANISM: Homo sapiens
FEATURE:
NAME/KEY: unsure
LOCATION: 7421
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 7427
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 11609
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 12605
OTHER INFORMATION: unknown
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LOCATION: 12742
OTHER INFORMATION: unknown
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OTHER INFORMATION: unknown
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LOCATION: 29422
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 29979
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 29980
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 29981
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 30136
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NAME/KEY: unsure
LOCATION: 30140
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 31205
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NAME/KEY: unsure
LOCATION: 31206
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 31592
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 33095
OTHER INFORMATION: unknown

NAME/KEY: unsure
LOCATION: 33160
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 34066
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 34072
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 36816
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 39020
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 42164
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 42459
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 46808
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OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 63290
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 66614
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 68660
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 68697
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 68718
OTHER INFORMATION: unknown
NAME/KEY: unsure
LOCATION: 68733
OTHER INFORMATION: unknown
NAME/KEY: unsure

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/ LOCATION: 68739
/ OTHER INFORMATION: unknown
/ NAME/KEY: unsure
/ LOCATION: 69785
/ OTHER INFORMATION: unknown
/ NAME/KEY: unsure
/ LOCATION: 79134
/ OTHER INFORMATION: unknown
/ NAME/KEY: unsure
/ LOCATION: 79198
/ OTHER INFORMATION: unknown
/ NAME/KEY: unsure
/ LOCATION: 8636
/ OTHER INFORMATION: unknown
/ OTHER INFORMATION:
US-09-791-211-3

Query Match
Best Local Similarity 4.7%; Score 47; DB 4; Length 87543;
Matches 56; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 922 GTATTGGAGTTCATGCAAAATGAGTGTGTTTAGCTGCTTGGCCACAAAAA 981
Db 19087 GTATTGGAGTTCGCTTAATGAGTGGATTTCAGCTGCTTTGCCACAAAA 19028

QY 982 AAAAAA 992
Db 19027 ATGGTTAACAA 19017

RESULT 5
US-09-007-005-17
; Sequence 17, Application US/09007005B
; Patent No. 6258558
; GENERAL INFORMATION:
; APPLICANT: Szostak, Jack W.
; APPLICANT: Roberts, Richard W.
; APPLICANT: Liu, Rihe
; TITLE OF INVENTION: SELECTION OF PROTEINS USING RNA-PROTEIN
; FILE REFERENCE: 00786/350003
; CURRENT APPLICATION NUMBER: US/09/007,005B
; CURRENT FILING DATE: 1998-01-14
; EARLIER APPLICATION NUMBER: 60/035,963
; EARLIER FILING DATE: 1997-01-27
; EARLIER APPLICATION NUMBER: 60/064,491
; EARLIER FILING DATE: 1997-11-06
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17
; LENGTH: 289
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Translation template
; NAME/KEY: misc feature
; LOCATION: (1)...(289)
; OTHER INFORMATION: n = A,T,C or G
US-09-007-005-17

Query Match
Best Local Similarity 4.4%; Score 43.6; DB 3; Length 289;
Matches 27; Conservative 85; Mismatches 111; Indels 0; Gaps 0;

QY 769 ACAGGACGCCCGCTGAGTGAAGAGGAGTTGGGGGTTTCAGATAGGATCGGG 828
Db 65 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 124
QY 829 AGGTGAGGACGCAAGACGACCCATGTAGATGAACCGTCAGAGACCCAGCACGG 888
Db 125 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 184
QY 889 CAGGAGCTCAGCGCCATCAGCGTCTGCTTATTTGGAGTTTCATGAAATGAGTG 948
Db 185 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 244
QY 949 TGTTTAGCTGCTTGGCCACAAAAA 991
Db 245 RCRURGRGRCRUAAAAA 287

RESULT 6
US-09-244-796-17
; Sequence 17, Application US/09244796
; Patent No. 6281344
; GENERAL INFORMATION:
; APPLICANT: Szostak, Jack W.
; APPLICANT: Roberts, Richard W.
; APPLICANT: Liu, Rihe
; TITLE OF INVENTION: SELECTION OF PROTEINS USING RNA-PROTEIN
; FILE REFERENCE: 00786/350007
; CURRENT APPLICATION NUMBER: US/09/244,796
; CURRENT FILING DATE: 1999-02-05
; EARLIER APPLICATION NUMBER: 60/035,963
; EARLIER FILING DATE: 1997-01-27
; EARLIER APPLICATION NUMBER: 60/064,491
; EARLIER FILING DATE: 1997-11-06
; EARLIER APPLICATION NUMBER: 09/007,005
; EARLIER FILING DATE: 1998-01-14
; NUMBER OF SEQ ID NOS: 33
; SOFTWARE: FastSeq for Windows Version 4.0
; SEQ ID NO 17
; LENGTH: 289
; TYPE: RNA
; ORGANISM: Artificial Sequence
; FEATURE:
; OTHER INFORMATION: Translation template
; NAME/KEY: misc feature
; LOCATION: (1)...(289)
; OTHER INFORMATION: n = A,T,C or G
US-09-244-796-17

Query Match
Best Local Similarity 4.4%; Score 43.6; DB 3; Length 289;
Matches 27; Conservative 85; Mismatches 111; Indels 0; Gaps 0;

QY 769 ACAGGACGCCCGCTGAGTGAAGAGGAGTTGGGGGTTTCAGATAGGATCGGG 828
Db 65 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 124
QY 829 AGGTGAGGACGCAAGACGACCCATGTAGATGAACCGTCAGAGACCCAGCACGG 888
Db 125 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 184
QY 889 CAGGAGCTCAGCGCCATCAGCGTCTGCTTATTTGGAGTTTCATGAAATGAGTG 948
Db 185 RNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRNRN 244
QY 949 TGTTTAGCTGCTTGGCCACAAAAA 991
Db 245 RCRURGRGRCRUAAAAA 287

RESULT 7
US-08-232-463-14/c
; Sequence 14, Application US/08232463
; Patent No. 5670367
; GENERAL INFORMATION:
; APPLICANT: DORNER, F.
; APPLICANT: SCHEIFLINGER, F.
; APPLICANT: FALKNER, F. G.
; TITLE OF INVENTION: RECOMBINANT FOWLPOX VIRUS
; NUMBER OF SEQUENCES: 52
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; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Foley & Lardner
; STREET: 1800 Diagonal Road, Suite 500
; CITY: Alexandria
; STATE: VA
; COUNTRY: USA
; ZIP: 22313-0299
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: PatentIn Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/232,463
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US/07/935,313
; FILING DATE:
; APPLICATION NUMBER: EP 91 114 300.6
; FILING DATE: 26-AUG-1991
; ATTORNEY/AGENT INFORMATION:
; NAME: BENT, Stephen A.
; REGISTRATION NUMBER: 29,768
; REFERENCE/DOCKET NUMBER: 30472/114 IMMU
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (703) 836-9300
; TELEFAX: (703) 683-4109
; TELEEX: 899149
; INFORMATION FOR SEQ ID NO: 14:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 7218 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
; IMMEDIATE SOURCE:
; CLONE: pTZgpt-Fls
US-08-232-463-14

Query Match 4.2%; Score 41.4; DB 1; Length 7218;
Best Local Similarity 1.7%; Pred. No. 0.48;
Matches 6; Conservative 203; Mismatches 144; Indels 0; Gaps 0;
QY 640 TCTGTGTCAGGAGTCAACAGCCCAATGTCACGACAGAGCCCTCACAGTGTGCCCC 699
Db 1442 TTGGTACRNR 1383
QY 700 CAGGTGTCAGCAGAGAGTGGAGGACTGGCAGGTCCTCCCTGGAGAGCCCATCTTGCT 759
Db 1382 RRRNR 1323
QY 760 TCCCGCTTACAGGAGCACCCTGCTGAGTGAAGAGGATTTGGGGGTTTCAAGATAG 819
Db 1322 RRRNR 1263
QY 820 GGAATGGGAGTCAAGAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC 879
Db 1262 RRRNR 1203
QY 880 CAAGCAGGTCAGGAGTCAAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGCAGC 939
Db 1202 RRRNR 1143
QY 940 AATAGTGTGTTTGTAGTCTCTTGGCCACAAAAAAGAAAAAAGAAAAA 992
Db 1142 RRRNR 1090

RESULT 8
US-09-252-991A-11640/c
; Sequence 11640, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:

; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11640
; LENGTH: 480
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11640

Query Match 4.1%; Score 40.6; DB 4; Length 480;
Best Local Similarity 44.4%; Pred. No. 0.29;
Matches 163; Conservative 0; Mismatches 204; Indels 0; Gaps 0;
QY 260 GAACATCAGGTGGCCCAAGAGGTGGTGAAGACCCACGAGCGGCTCTTCAACCTCA 319
Db 424 GGAGATCAAGAGAGCATTGCGAGAGGTGGTGGCCAGCGCTGATCTACCTGCCCTCCG 365
QY 320 CGTCACTCAAGTCCAGTCCAGACTGCTCACTTCTTCTGCGGGGCTCTTCCACCTC 379
Db 364 CGTTCGGACCTGACAACTCCGCACTCGACAAGTACCTTCCACTATTGCGGGGCTC 305
QY 380 AGTGCCCATGTGGACAGTGGCCAGGTCAGAGTACAGTACAGTGGAGTGTGTCAAGCAGT 439
Db 304 CGCGCGCATGGGCCACCGGAGCGGATCAAGATCTCAAGTGTCTCTGGAGCCATCGG 245
QY 440 GTCTGAGCTGGGGCCAACTTCACTCTGACAGACAGAGGGGAGCCGCCAGGTGGAGT 499
Db 244 CAGCAGTTTCGGCGGCGCCACGAGTGTACAGATCAACTAGCGCGGAGCAGGACGA 185
QY 500 GATTCGCCAGGCTCTCTGGGCGAGCCCACTATCAACCAAGCTGATCGGAAGATGG 559
Db 184 GATCGCATGAGGCGCTTGGCCAGCGGTTCGGCAGCGGGCGATGAAGGGCATGCTCG 125
QY 560 GCAGTCCACTGTCAGAGAGACCATGTCACAGGAGCGCTGCAACTTCTCTTCTGCTGCC 619
Db 124 CATGTCGAGCAGTGCATGGGCGACTAGCAGAGAACGCTGGACCGTGGCCGACGCTGCA 65
QY 620 GAGCCAG 626
Db 64 CAACCCG 58

RESULT 9
US-09-252-991A-11799
; Sequence 11799, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11799
; LENGTH: 1575
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11799
Query Match 4.1%; Score 40.6; DB 4; Length 1575;

Best Local Similarity 44.4%; Pred. No. 0.44;
Matches 163; Conservative 0; Mismatches 204; Indels 0; Gaps 0;

QY	260	GAACATCAAGTGGCCAAAGAGGTGGTGAAGACCAACAGAGCCGGCCTCTTTCAACCTCAA	319
Db	1173	GGAGATCAAGAAGACCATCGAGCAGGTGGTCGCCAGGGCCTGATCTACCTGGCCTCGG	1232
QY	320	CGTCACACTCAAGTCCAGTCCAGACCTGCTCACTACTTCTGCGGGGCTCCTCCACCTC	379
Db	1233	CGTTCCGGACCTGCACAATCCGCAACTCGACAAGTACTCTTCCACCTATTTCGCGGGTCT	1292
QY	380	AGGTGCCCATCTGGACAGTGCAGGCTACAGATGCATCTGGGAGCTGTGTTCCAGACCAGT	439
Db	1293	CGGCGGCATCGGCCACCGGGAGCGGATCAAGATCCTCAAGCTGCTTGGGAGCGCATCGG	1352
QY	440	GTCGTAGCTGGCGCCAACTTCACTCTCAGACAGAGGGCAGGCCCTCCAGGTGGAGAT	499
Db	1353	CAGCGATTTCGGCGGCCGCCACGAGCTGTACGAGATCAACTACGCCGGCAGCAGGACGA	1412
QY	500	GATCTGCCAGGCTCTCGGGCAGCCACCTATTCACCAACAGCTGTATTCGGGAAGGATGG	559
Db	1413	GATCCGGCATCGAGGCCCTTGGCCCAAGGCGGTTCGGCAGGGGGCGGATGAAGGCGCATCTCGG	1472
QY	560	GCAGGTCCACTGCAGCAGAGACCATGCCACAGGCAGCCTTCGCCAACTTCTTCCTCTCGCC	619
Db	1473	CATGGTTCAGCATGTGATGGCGCACTACGACGAGACGGGTGGACCGTGGCGACCTTGCA	1532
QY	620	GAGCCAG	626
Db	1533	CAACCCG	1539

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RESULT 10
US-09-252-991A-11716
; Sequence 11716, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS

```

Query Match	4.1%	Score 40.6;	DB 4;	Length 2022;
Best Local Similarity	44.4%;	Pred. No. 0.48;		
Matches 163; Conservative		0; Mismatches 204;	Indels 0;	Gaps 0;

QY	260	GAACATCAAGGTGGCCAAAGAGTGGTGAAGACCCACGAGCGCGCTCTTCAACCTCAA	319
Db	1078	GGAGATCAAGAACCATTGAGCAGGTGGTCGCCAGCGGCTGATCTACCTGCCCTCGG	1137
QY	320	CGTCACACTCAAGTCCAGTCCAGACTGCTCACTACTTTCGCGGGGTCCTCCACCTC	379
Db	1138	CGTTGGGACCTGCACAAATCCGACACTCGACAAGTACTCTTCCACCTATTTCGCGGCTC	1197
QY	380	AGGTGCCCATGTGGACAGTCCCAAGGCTACAGATGCACCTGGGAGGTGGTCTCAAGCCAGT	439
Db	1198	CGGCGGCATGGGCCACCGGGAGCGGATCAAGATCCTCAAGTCTCTGGGAGGCCATCGG	1257
QY	440	GTCTGAGCTCGGGCCACTTCACTCTGACGAGCAGAGGGGCGAGGCCCGCAGGGTGGAGAT	499
Db	1258	CAGCGAGTTTCGGCGGCGCCACAGAGTGTACAGATCAACTACCGCGGACGACAGACGA	1317

QY	500	GATCTGCAGCGCTCTCGGCGCCGCCACCTATCACCACAGCCTTGATCGGGAAGATGG	559
Db	1318	GATCGCATGCGGCGCTCTGCCCGAGGCGGTGGGAGCGGGCGATGAAGGGCATGCTCGG	1377
QY	560	GCAGGTCACTGTGCAGCAGAGACCATGCCACAGGCAGCCTGCCAACTTCTCTCTCTCTCC	619
Db	1378	CATGGTCGAGCAGTGCATGGGCGCACTACGACGAGAAAGGCTGGACCGTGGCGCACCTGCA	1437
QY	620	GAGCCAG	626
Db	1438	CAACCCG	1444

RESULT 11
 US-09-252-991A-10993
 ; Sequence 10993, Application US/09252991A
 ; Patent No. 6551795
 ; GENERAL INFORMATION:
 ; APPLICANT: Marc J. Rubenfield et al.
 ; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
 ; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
 ; FILE REFERENCE: 107196.136
 ; CURRENT APPLICATION NUMBER: US/09/252,991A
 ; CURRENT FILING DATE: 1999-02-18
 ; PRIOR APPLICATION NUMBER: US 60/074,788
 ; PRIOR FILING DATE: 1998-02-18
 ; PRIOR APPLICATION NUMBER: US 60/094,190
 ; PRIOR FILING DATE: 1998-07-27
 ; NUMBER OF SEQ ID NOS: 33142
 ; SEQ ID NO 10993
 ; LENGTH: 1332
 ; TYPE: DNA
 ; ORGANISM: Pseudomonas aeruginosa
 US-09-252-991A-10993

Query Match 4.1%; Score 40.2; DB 4; Length 1332;
Best Local Similarity 49.8%; Pred. No. 0.53;
Matches 102; Conservative 0; Mismatches 103; Indels 0; Gaps 0;

QY	218	CGAGCACCACCCCCATCACTATTCCCTCTGTGGAAACCAAGAACATCAAGGTGGCCAA	277
Db	826	CCGCCACCCGACGAGGATCGCCGGATCGCGGGATGCGGAATACCACTCGGTGACCGA	885
QY	278	GAAGGTGGTGAACACCCACGAGCGGCGCTCTTCAACCTCAAGTCACTCAAGTCCAG	337
Db	886	GCCGATGGCGATGGTCGCGGTGACCGCCACCATCAGTCTGTAGCGACCATGACACCCC	945
QY	338	TCCAGACCTGCTCACCTACTTCTGCCGGGGCTCTCCACCTCAGGTGCCCATGTGGACAG	397
Db	946	GCGCATCAGCGCGGAGGCGCTGTGTAGCGCTGCTCCAGCACCTGGGTACGGTGAACAC	1005
QY	398	TGCCAGGCTACAGATGCATCTGGAG	422
Db	1006	CCGACGGCGCGGATCTGCGCGAG	1030

RESULT 12
 US-09-252-991A-11258/c
 ; Sequence 11258, Application US/09252991A
 ; Patent No. 6551795
 ; GENERAL INFORMATION:
 ; APPLICANT: Marc J. Rubenfield et al.
 ; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
 ; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
 ; FILE REFERENCE: 107196.136
 ; CURRENT APPLICATION NUMBER: US/09/252,991A
 ; CURRENT FILING DATE: 1999-02-18
 ; PRIOR APPLICATION NUMBER: US 60/074,788
 ; PRIOR FILING DATE: 1998-02-18
 ; PRIOR APPLICATION NUMBER: US 60/094,190
 ; PRIOR FILING DATE: 1998-07-27
 ; NUMBER OF SEQ ID NOS: 33142

```

; SEQ ID NO 11258
; LENGTH: 1656
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11258

Query Match      4.1%; Score 40.2; DB 4; Length 1656;
Best Local Similarity 49.8%; Pred. No. 0.57;
Matches 102; Conservative 0; Mismatches 103; Indels 0; Gaps 0;

QY 218 CCAGCCACCACCGCCCATCACTATTCCCTCTGTGGAACCAAGAACATCAAGGTGGCCAA 277
Db 570 CCCGCCACCGCAGAGGATCGCGGGATCCGCGGATCCGGAATACCACTCGGTGACCGA 511

QY 278 GAAGGTGGTGAAGACCCACGAGCGGGCTCTTCAAGCTCAACCTCACTCAAGTCCAG 337
Db 510 GCCGATGGCGATGTCGCGGTGACCGCCACCATCAGTTCGTAGGCGACCATGACCCCC 451

QY 338 TCAGACCTGCTCACTACTTCTTCGCGGGGTCTCTCACTCAGGTGCCCATGTGACAG 397
Db 450 GCCGATCAGCGCGGAGGAGCGCTGTGTAGCGTGTCTCAGCACCTGGGTACCGGTGAAC 391

QY 398 TGCAGGCTACAGATGCACCTGGGAG 422
Db 390 CCGCAGCGCGCGATCTGCCCGAG 366

RESULT 13
US-09-252-991A-11190/c
; Sequence 11190, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; PRIOR FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11190
; LENGTH: 2217
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
; FEATURE:
; NAME/KEY: unsure
; LOCATION: (2217)
; OTHER INFORMATION: Identity of nucleotide at the above locations are unknown.
US-09-252-991A-11190

Query Match      4.1%; Score 40.2; DB 4; Length 2217;
Best Local Similarity 49.8%; Pred. No. 0.63;
Matches 102; Conservative 0; Mismatches 103; Indels 0; Gaps 0;

QY 218 CCAGCCACCACCGCCCATCACTATTCCCTCTGTGGAACCAAGAACATCAAGGTGGCCAA 277
Db 778 CCCGCCACCGCAGAGGATCGCGGGATCCGCGGATCCGGAATACCACTCGGTGACCGA 719

QY 278 GAAGGTGGTGAAGACCCACGAGCGGGCTCTTCAAGCTCAACCTCACTCAAGTCCAG 337
Db 718 GCCGATGGCGATGTCGCGGTGACCGCCACCATCAGTTCGTAGGCGACCATGACCCCC 659

QY 338 TCAGACCTGCTCACTACTTCTTCGCGGGGTCTCTCACTCAGGTGCCCATGTGACAG 397
Db 658 GCGATCAGCGCGGAGAGCGCTGTGTAGCGTGTCTCAGCACCTGGGTACCGGTGAAC 599

QY 398 TGCAGGCTACAGATGCACCTGGGAG 422
Db 598 CCGCAGCGCGCGATCTGCCCGAG 574

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US-09-103-840A-1

Query Match 3.8%; Score 37.8; DB 3; Length 4411529;
Best Local Similarity 47.3%; Pred. No. 34;
Matches 114; Conservative 0; Mismatches 127; Indels 0; Gaps 0;

QY 338 TCAGACCTGCTCACCTACTTTCGCGGCGCTCTCCACCTCAGGTGCCATGTGGACAG 397
DB 284000 TTCCGACCTTGTGACATCTCGGCTCGCGGTACGGCGCGCGGACGGGATCCCGCCAC 283941
QY 398 TGCAGGCTACAGATCACTGGAGCTGTGGTCCAAAGCCAGTGTCTGAGCTCGGGGCCAA 457
DB 283940 CGCCTGGACCGGCGCGGCGGTGTGCAACACAAACCCCGGACGCTGACGCTGAC 283881
QY 458 CTTCACTCTGCAGGACAGAGGGGCGAGGCCCGGAGGTGGAGATGATCTGCAGGCGTCTTC 517
DB 283880 CTTGCGCGGCGCCACCGTGTGACCGGCGTGGCGCGCGCGGCGGCGGCGGCGGCGG 283821
QY 518 GGGACGCCACCTATCACCAACAGCTGATCGGGAAGGATGGGAGGTCCACCTGCAGCA 577
DB 283820 GGGCCATCGACGGTGTGGCCATCAACTTGGCGACGGCCCCCAGGTCCGACAACTGCA 283761
QY 578 G 578
DB 283760 G 283760

RESULT 16

US-09-522-714-21

; Sequence 21, Application US/09522714
; Patent No. 6563020
; GENERAL INFORMATION:
; APPLICANT: Simmons, Carl R.
; APPLICANT: Yalpani, Nasser
; TITLE OF INVENTION: Maize Chitinases and Their Use in
; FILE REFERENCE: 1100
; CURRENT APPLICATION NUMBER: US/09/522,714
; EARLIER FILING DATE: 2000-03-10
; EARLIER APPLICATION NUMBER: 60/125,915
; EARLIER FILING DATE: 1999-03-24
; NUMBER OF SEQ ID NOS: 32
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 21
; LENGTH: 1048
; TYPE: DNA
; ORGANISM: Zea mays
; FEATURE:
; NAME/KEY: CDS
; LOCATION: (42)...(854)
US-09-522-714-21

Query Match 3.8%; Score 37.6; DB 4; Length 1048;
Best Local Similarity 42.6%; Pred. No. 2.2;
Matches 196; Conservative 0; Mismatches 264; Indels 0; Gaps 0;

QY 83 CTGCTTGGCGGTGCTGCCAGAGCTTCTCCAGGACGCGGAGGAGAAATTAACCC 142
DB 326 CCGGACGCTTCTTGGCGCGCGCGGCTACTACCGGGGTTCGGCGCGGACCGGT 385
QY 143 TGTGTCTTCAATGCTTCAAGTCTCTGGAAGTTTTTCCCAAGGCGCGGTGCTCAT 202
DB 386 CGACGACTCCAGCGGAGATCGCGGCTTCTTGGCAACGCCACACGAGACATATA 445
QY 203 AACCTGCTGTGACCGGACGACACCGCCCATCTATCTCTCTGTGGACCAAGAA 262
DB 446 GTTCTGTACATCAACGAGATCGACGGGCGGAGAACTACTCGGACCGGAAACAC 505
QY 263 CATCAAGTGGCAAGAGGTGGTGAAGACCAACGCGGCGCTCTTCAACCTCAAGT 322
DB 506 GCAGTGGCGGTGCCAGGCGGGAAGGGTACTACGGCGGCGGCGGCGGCGGCTG 565
QY 323 CACACTCAAGTCCAGTCAGACCTGCTCACTTCTCTCGGCGGCGCTCTCCACCTCAG 382

DB 566 GAACCTTCAACTACGGGCGCGCGGCGGCGAGAGCATCGGCTTCGACGGGCTGGGCGACCCCGA 625
QY 383 TGCCCATGTGGACAGTGCAGGCTACAGATGACACTGGGAGCTGTGGTCCAGCCAGTGTC 442
DB 626 CGCGGTGGCGGCGAGCGCGGTCTCGGCTTCGGCTCCGGCTCTGTGTAAGTGAACA 685
QY 443 TGAGTGGCGGCGCAACTTCATCTTCAGGACAGAGGGGCGAGCCCGGAGGATGAT 502
DB 686 CGTGACGCGGCGCATCGTCTCGGCGAGGCTTCGGCGCCACCATCGGGCGCATCAACGG 745
QY 503 CTGCCAGGCTCTCGGCGAGCGCCACCTATCACCAACAGC 542
DB 746 CGCGTCTGAGTGGAGCGGCAAGAACCCCACTCCGCTCAAC 785

RESULT 17

US-08-123-161A-11
; Sequence 11, Application US/08123161A
; Patent No. 5449616

; GENERAL INFORMATION:
; APPLICANT: Campbell, Kevin P.
; APPLICANT: Roberts, Steven L.
; APPLICANT: Anderson, Richard D.
; APPLICANT: Ibraghimov, Oxana B.
; APPLICANT: Yang, Bin
; TITLE OF INVENTION: NUCLEIC ACID ENCODING DYSTROPHIN-ASSOCIATED
; NUMBER OF SEQUENCES: 14
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kevin M. Farrell, P.C.
; STREET: P.O. Box 999
; CITY: York Harbor
; STATE: ME
; COUNTRY: USA
; ZIP: 03911
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/123,161A
; FILING DATE: 16-SEP-93
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 07/946,234
; FILING DATE: 14-SEP-92
; ATTORNEY/AGENT INFORMATION:
; NAME: Farrell, Kevin M.
; REGISTRATION NUMBER: 35,505
; REFERENCE/DOCKET NUMBER: UIRF89-11A4
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (207) 363-0558
; TELEFAX: (207) 363-0528
; INFORMATION FOR SEQ ID NO. 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1396 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 4..1164
US-08-123-161A-11

Query Match 3.8%; Score 37.6; DB 1; Length 1396;
Best Local Similarity 65.5%; Pred. No. 2.5;
Matches 55; Conservative 0; Mismatches 29; Indels 0; Gaps 0;

QY 909 GCGTGCACCTGTCTGTTATTTGGAGTTTCATGCAAAATAGTGTGTTTACGTGCTCTGCCA 968

Db 1309 GGGTGGGTCAGACTGTCTGGAGTAAGACATTTCAGATAAATATCTGCTCTGCTCA 1368
QY 969 CAAAAAAAAAAAAAAAAAAAAA 992
Db 1369 CAAAAAAAAAAAAAAAAAAAAA 1392

RESULT 18
US-08-483-278-11
; Sequence 11, Application US/08483278
; Patent No. 5686073
; GENERAL INFORMATION:
; APPLICANT: Campbell, Kevin P.
; APPLICANT: Ibrahimov, Oxana B.
; APPLICANT: Ervasti, James M.
; APPLICANT: Leveille, Cynthia J.
; TITLE OF INVENTION: NUCLEIC ACID ENCODING DYSTROPHIN-ASSOCIATED
; TITLE OF INVENTION: PROTEIN
; NUMBER OF SEQUENCES: 15
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Kevin M. Farrell, P.C.
; STREET: P.O. Box 999
; CITY: York Harbor
; STATE: ME
; COUNTRY: USA
; ZIP: 03911
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
; OPERATING SYSTEM: PC-DOS/MS-DOS
; SOFTWARE: Patent In Release #1.0, Version #1.25
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/483,278
; FILING DATE:
; CLASSIFICATION: 435
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: 08/123,161
; FILING DATE: 16-SEP-93
; ATTORNEY/AGENT INFORMATION:
; NAME: Farrell, Kevin M.
; REGISTRATION NUMBER: 35,505
; REFERENCE/DOCKET NUMBER: UIRP89-11A5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: (207) 363-0558
; TELEFAX: (207) 363-0528
; INFORMATION FOR SEQ ID NO: 11:
; SEQUENCE CHARACTERISTICS:
; LENGTH: 1396 base pairs
; TYPE: nucleic acid
; STRANDEDNESS: double
; TOPOLOGY: linear
; MOLECULE TYPE: cDNA
; FEATURE:
; NAME/KEY: CDS
; LOCATION: 4...1164

US-08-483-278-11
Query Match 3.8%; Score 37.6; DB 1; Length 1396;
Best Local Similarity 65.5%; Pred. No. 2.5;
Matches 55; Conservative 0; Mismatches 29; Indels 0; Gaps 0;
QY 909 GCGTGCACTGTTGCTATTTGGAGTTCATGCAAAATGAGTGTCTTTTAGCTGCTCTTGCCA 968
Db 1309 GGGTGGGTCAGAGTGTGGAGTTCAGATAAATATCTGCTCTGCTCA 1368
QY 969 CAAAAAAAAAAAAAAAAAAAAA 992
Db 1369 CAAAAAAAAAAAAAAAAAAAAA 1392

RESULT 19
US-09-252-991A-5567/c
; Sequence 5567, Application US/09252991A

; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 5567
; LENGTH: 1992
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-5567

Query Match 3.8%; Score 37.6; DB 4; Length 1992;
Best Local Similarity 50.6%; Pred. No. 2.8;
Matches 91; Conservative 0; Mismatches 89; Indels 0; Gaps 0;
QY 515 CTCGGGCAGCCACCTATCACCAACAGCTGATCGGGAAGGATGGCGAGGTCCACCTGCA 574
Db 1569 CTCGTGCGCGCGCGCGCTCCATGAACAGAGATCAGGGCTGCGGTAGAGCGGTAGAA 1510
QY 575 GCAGAGACCATGCCACAGCAGCGCTGCCAACTTCTCTTCTCCGCGAGCCAGACATCGGA 634
Db 1509 GAGCAGTCCGGCAGCAGAGCTCCAGCGGCCCATCTCTGCGGGCCGACGGCTTCGTGAT 1450
QY 635 CTGGTTCTGTCGTCAGGCTGCAAAACAAACCCCAATGTCACAGCAGCGCCCTCACAGTGGT 694
Db 1449 CTGCTGCTGAGCAGGCTTTCGAGGAATATCCAGGCGCAGGCGGACAGGCGTGGT 1390

RESULT 20
US-08-997-080-179/c
; Sequence 179, Application US/08997080
; Patent No. 5968524
; GENERAL INFORMATION:
; APPLICANT: WATSON, JAMES D.
; APPLICANT: TAN, PAUL L.J.
; TITLE OF INVENTION: METHODS AND COMPOUNDS FOR THE TREATMENT OF IMMUNOLOGICALLY-
; NUMBER OF SEQUENCES: 194
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Law Offices of Ann W. Speckman
; STREET: 2601 Elliott Avenue, Suite 4185
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98121
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/997,080
; FILING DATE:
; CLASSIFICATION:
; PRIOR APPLICATION DATA:
; APPLICATION NUMBER:
; FILING DATE:
; ATTORNEY/AGENT INFORMATION:
; NAME: Sleath, Janet
; REGISTRATION NUMBER: 37,007
; REFERENCE/DOCKET NUMBER: 11000.1007
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 206-269-0565
; TELEFAX: 206-269-0563
; TELEX:
; INFORMATION FOR SEQ ID NO: 179:

```
/
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 520 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/
US-08-997-080-179

Query Match          3.8%; Score 37.4; DB 2; Length 520;
Best Local Similarity 48.0%; Pred. No. 1.9;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 374 CACCTCAGTGCCTCCATGTGACAGTGCAGGCTACAGATGCACTGGAGCTGTGTCCAA 433
DB 324 CTCCTCGGTGAGCTCGGGATCGACACCGGATGATGTTGCCGTGCGGATTGAC 265
QY 434 GCCAGTGTCTGAGCTGCGGGCCAACTTCACTCTCAGGACAGAGGGGCGAGCCCGAGGGT 493
DB 264 GCCAGGTTCGGAGTTGCGGATCGCATCTCGATGAGGCGCAGCTGGCTCGCTCGTAGGG 205
QY 494 GGAGATGATCTGCCAGGCGTCTCTCGGGCAGCCCACTATCACCACAGCCTGTATCGGGAA 553
DB 204 CTTGATCACCACCATGCGCGCTCGGGCAGCTGATGCTGGACAGCTCGTGATCGGGGT 145
QY 554 GGATGGCGAGTCCACCTGCGAGCAGAGACCATGCCACAGGCGAG 596
DB 144 GGAGCGCGGTAGTAGTGTGATCGGTTGACATGCGG 102

RESULT 21
US-08-997-362-179/c
/ Sequence 179, Application US/08997362
/ Patent No. 5985287
/ GENERAL INFORMATION:
/ APPLICANT: Tan, Paul
/ APPLICANT: Hiyama, Jun
/ APPLICANT: Visser, Elizabeth
/ APPLICANT: Skinner, Margot
/ APPLICANT: Scott, Linda
/ APPLICANT: Prestidge, Ross
/ TITLE OF INVENTION: COMPOUNDS AND METHODS FOR
/ TITLE OF INVENTION: TREATMENT AND DIAGNOSIS OF MYCOBACTERIAL INFECTIONS
/ NUMBER OF SEQUENCES: 194
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: Law Offices of Ann W. Speckman
/ STREET: 2601 Elliott Avenue, Suite 4185
/ CITY: Seattle
/ STATE: WA
/ COUNTRY: USA
/ ZIP: 98121
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Diskette
/ COMPUTER: IBM Compatible
/ OPERATING SYSTEM: DOS
/ SOFTWARE: FastSeq for Windows Version 2.0
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/08/997,362
/ FILING DATE:
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: U.S. Patent Application No. 5985287 08/873,970
/ FILING DATE: June 12, 1997
/ APPLICATION NUMBER: U.S. Patent Application No. 5985287 08/705,347
/ FILING DATE: August 29, 1996
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Sleath, Janet
/ REGISTRATION NUMBER: 37,007
/ REFERENCE/DOCKET NUMBER: 11000.1002c2
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 206-269-0565
/ TELEFAX: 206-269-0563
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 179:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 520 base pairs
```

```
/
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 520 base pairs
/ TYPE: nucleic acid
/ STRANDEDNESS: single
/ TOPOLOGY: linear
/
US-08-997-362-179

Query Match          3.8%; Score 37.4; DB 2; Length 520;
Best Local Similarity 48.0%; Pred. No. 1.9;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 374 CACCTCAGTGCCTCCATGTGACAGTGCAGGCTACAGATGCACTGGAGCTGTGTCCAA 433
DB 324 CTCCTCGGTGAGCTCGGGATCGACACCGGATGATGTTGCCGTGCGGATTGAC 265
QY 434 GCCAGTGTCTGAGCTGCGGGCCAACTTCACTCTCAGGACAGAGGGGCGAGCCCGAGGGT 493
DB 264 GCCAGGTTCGGAGTTGCGGATCGCATCTCGATGAGGCGCAGCTGGCTCGCTCGTAGGG 205
QY 494 GGAGATGATCTGCCAGGCGTCTCTCGGGCAGCCCACTATCACCACAGCCTGTATCGGGAA 553
DB 204 CTTGATCACCACCATGCGCGCTCGGGCAGCTGATGCTGGACAGCTCGTGATCGGGGT 145
QY 554 GGATGGCGAGTCCACCTGCGAGCAGAGACCATGCCACAGGCGAG 596
DB 144 GGAGCGCGGTAGTAGTGTGATCGGTTGACATGCGG 102

RESULT 22
US-09-095-855-179/c
/ Sequence 179, Application US/09095855
/ Patent No. 6160093
/ GENERAL INFORMATION:
/ APPLICANT: Tan, Paul
/ APPLICANT: Visser, Elizabeth
/ APPLICANT: Skinner, Margot
/ APPLICANT: Prestidge, Ross
/ TITLE OF INVENTION: Compounds and Methods for
/ TITLE OF INVENTION: Treatment and Diagnosis of Mycobacterial Infections
/ NUMBER OF SEQUENCES: 208
/ CORRESPONDENCE ADDRESS:
/ ADDRESSEE: Law Offices of Ann W. Speckman
/ STREET: 2601 Elliott Avenue, Suite 4185
/ CITY: Seattle
/ STATE: WA
/ COUNTRY: USA
/ ZIP: 98121
/ COMPUTER READABLE FORM:
/ MEDIUM TYPE: Diskette
/ COMPUTER: IBM Compatible
/ OPERATING SYSTEM: DOS
/ SOFTWARE: FastSeq for Windows Version 2.0
/ CURRENT APPLICATION DATA:
/ APPLICATION NUMBER: US/09/095,855
/ FILING DATE:
/ CLASSIFICATION:
/ PRIOR APPLICATION DATA:
/ APPLICATION NUMBER: 08/705,347
/ FILING DATE: 29-AUG-1996
/ APPLICATION NUMBER: 08/873,970
/ FILING DATE: 12-JUN-1997
/ APPLICATION NUMBER: 08/997,362
/ FILING DATE: 23-DEC-1997
/ ATTORNEY/AGENT INFORMATION:
/ NAME: Sleath, Janet
/ REGISTRATION NUMBER: 37,007
/ REFERENCE/DOCKET NUMBER: 11000.1002c3
/ TELECOMMUNICATION INFORMATION:
/ TELEPHONE: 206-269-0565
/ TELEFAX: 206-269-0563
/ TELEX:
/ INFORMATION FOR SEQ ID NO: 179:
/ SEQUENCE CHARACTERISTICS:
/ LENGTH: 520 base pairs
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```
; TYPE: nucleic acid
; STRANDEDNESS: single
; TOPOLOGY: linear
US-09-095-855-179

Query Match      3.8%; Score 37.4; DB 3; Length 520;
Best Local Similarity 48.0%; Pred. No. 1.9;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 374 CACCTCAGGTGCCATGTGGACAGTGCAGGCTACAGATGCACCTGGAGCTGTGTCCTCAA 433
Db 324 CTCTCGGTGAGTGGGATGACACCCCGATGATGTTCCGCTGCTGTCGATGAC 265
QY 434 GCCAGTGTCTGAGTGGCGGCCCACTTCACTCTGCAGGACAGAGGGGCGAGGCCCGAGGGT 493
Db 264 GCCAGTGTGGAGTGGCGATGCATCTCGATGAGGCGAGCTGCCTCGCTAGGG 205
QY 494 GGAGATGATCTGCAGGCGTCTCGGCGAGCCACCTATCACCACAGCCTGATCGGGAA 553
Db 204 CTTGATCACCACCATCGCGCTCGGCGACGTTGATGCTGGACAGCTGCGTGTGATCGGGT 145
QY 554 GGATGGCGAGTCCACCTCGCAGCAGACCATGCCACAGGCAG 596
Db 144 GGAGGCGCCGTAGTGTGATGTTGATCCGGTTGAACATGCCG 102

RESULT 23
US-09-324-542-179/c
; Sequence 179, Application US/09324542
; Patent No. 6328978
; GENERAL INFORMATION:
; APPLICANT: Watson, James D.
; APPLICANT: Tan, Paul L.J.
; TITLE OF INVENTION: Methods and Compounds for the Treatment
; TITLE OF INVENTION: Of Immunologically-Mediated Skin Disorders
; FILE REFERENCE: 11000.1007c1
; CURRENT APPLICATION NUMBER: US/09/324,542
; EARLIER FILING DATE: 1999-06-02
; EARLIER APPLICATION NUMBER: US 08/997,080
; EARLIER FILING DATE: 1997-12-23
; NUMBER OF SEQ ID NOS: 194
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 179
; LENGTH: 520
; TYPE: DNA
; ORGANISM: Mycobacterium vaccae
US-09-324-542-179

Query Match      3.8%; Score 37.4; DB 4; Length 520;
Best Local Similarity 48.0%; Pred. No. 1.9;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 374 CACCTCAGGTGCCATGTGGACAGTGCAGGCTACAGATGCACCTGGAGCTGTGTCCTCAA 433
Db 324 CTCTCGGTGAGTGGGATGACACCCCGATGATGTTCCGCTGCTGTCGATGAC 265
QY 434 GCCAGTGTCTGAGTGGCGGCCCACTTCACTCTGCAGGACAGAGGGGCGAGGCCCGAGGGT 493
Db 264 GCCAGTGTGGAGTGGCGATGCATCTCGATGAGGCGAGCTGCCTCGCTAGGG 205
QY 494 GGAGATGATCTGCAGGCGTCTCGGCGAGCCACCTATCACCACAGCCTGATCGGGAA 553
Db 204 CTTGATCACCACCATCGCGCTCGGCGACGTTGATGCTGGACAGCTGCGTGTGATCGGGT 145
QY 554 GGATGGCGAGTCCACCTCGCAGCAGACCATGCCACAGGCAG 596
Db 144 GGAGGCGCCGTAGTGTGATGTTGATCCGGTTGAACATGCCG 102

RESULT 24
US-09-205-426-179/c
; Sequence 179, Application US/09205426
; Patent No. 6406704
; GENERAL INFORMATION:
; APPLICANT: Watson, James D.
; APPLICANT: Tan, Paul L. J.
; TITLE OF INVENTION: Compounds and Methods for Treatment and
; TITLE OF INVENTION: Diagnosis of Mycobacterial Infections
; FILE REFERENCE: 11000.1002c4
; CURRENT APPLICATION NUMBER: US/09/205,426
; CURRENT FILING DATE: 1998-12-04
; EARLIER APPLICATION NUMBER: 09/095,855
; EARLIER FILING DATE: 1998-06-11
; EARLIER APPLICATION NUMBER: 08/997,362
; EARLIER FILING DATE: 1997-12-23
; EARLIER APPLICATION NUMBER: 08/873,970
; EARLIER FILING DATE: 1997-06-12
; EARLIER APPLICATION NUMBER: 08/705,347
; EARLIER FILING DATE: 1996-08-23
; NUMBER OF SEQ ID NOS: 208
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 179
; LENGTH: 520
; TYPE: DNA
; ORGANISM: Mycobacterium vaccae
US-09-205-426-179

Query Match      3.8%; Score 37.4; DB 4; Length 520;
Best Local Similarity 48.0%; Pred. No. 1.9;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;

QY 374 CACCTCAGGTGCCATGTGGACAGTGCAGGCTACAGATGCACCTGGAGCTGTGTCCTCAA 433
Db 324 CTCTCGGTGAGTGGGATGACACCCCGATGATGTTCCGCTGCTGTCGATGAC 265
QY 434 GCCAGTGTCTGAGTGGCGGCCCACTTCACTCTGCAGGACAGAGGGGCGAGGCCCGAGGGT 493
Db 264 GCCAGTGTGGAGTGGCGATGCATCTCGATGAGGCGAGCTGCCTCGCTAGGG 205
QY 494 GGAGATGATCTGCAGGCGTCTCGGCGAGCCACCTATCACCACAGCCTGATCGGGAA 553
Db 204 CTTGATCACCACCATCGCGCTCGGCGACGTTGATGCTGGACAGCTGCGTGTGATCGGGT 145
QY 554 GGATGGCGAGTCCACCTCGCAGCAGACCATGCCACAGGCAG 596
Db 144 GGAGGCGCCGTAGTGTGATGTTGATCCGGTTGAACATGCCG 102

RESULT 25
US-09-095-855-202/c
; Sequence 202, Application US/09095855
; Patent No. 6160093
; GENERAL INFORMATION:
; APPLICANT: Tan, Paul
; APPLICANT: Visser, Elizabeth
; APPLICANT: Skinner, Margot
; APPLICANT: Prestidge, Ross
; TITLE OF INVENTION: Compounds and Methods for
; TITLE OF INVENTION: Treatment and Diagnosis of Mycobacterial Infections
; NUMBER OF SEQUENCES: 208
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Law Offices of Ann W. Speckman
; STREET: 2601 Elliott Avenue, Suite 4185
; CITY: Seattle
; STATE: WA
; COUNTRY: USA
; ZIP: 98121
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Diskette
; COMPUTER: IBM Compatible
; OPERATING SYSTEM: DOS
; SOFTWARE: FastSeq for Windows Version 2.0
; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/09/095,855
; FILING DATE:
```

CLASSIFICATION:
PRIOR APPLICATION DATA:
APPLICATION NUMBER: 08/705,347
FILING DATE: 29-AUG-1996
APPLICATION NUMBER: 08/873,970
FILING DATE: 12-JUN-1997
APPLICATION NUMBER: 08/997,362
FILING DATE: 23-DEC-1997
ATTORNEY/AGENT INFORMATION:
NAME: Sleath, Janet
REGISTRATION NUMBER: 37,007
REFERENCE/DOCKET NUMBER: 11000.1002c3
TELECOMMUNICATION INFORMATION:
TELEPHONE: 206-269-0565
TELEFAX: 206-269-0563
TELEX:
INFORMATION FOR SEQ ID NO: 202:
SEQUENCE CHARACTERISTICS:
LENGTH: 570 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: cDNA
US-09-095-855-202

Query Match 3.8%; Score 37.4; DB 3; Length 570;
Best Local Similarity 48.0%; Pred. No. 2;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;
QY 374 CACCTCAGTCCCATGTGGACAGTCCAGGCTACAGATGACCTGGGAGCTGTGTCCTCAA 433
Db 333 CTCCTCGGTGAGCTGCGGGATCGACACCCGATGATGTTGCCGTGCTGCGATTGAC 274
QY 434 GCCAGTGTCTGAGCTGCGGGCACTTCACTCTCAGACAGAGAGGGGAGGCTGATCGGAA 553
Db 273 GCCAGTGTCTGAGCTGCGGGATCGACACCCGATGATGTTGCCGTGCTGCGATTGAC 274
QY 494 GGAGATGATCTGCCAGGCTCTCTCGGGCAGCCACCTATCACCAACAGCTGATCGGAA 553
Db 213 CTTGATCACCACCATGCGGCTCGGGCAGCTTGTGATCCGGTTGACATGCCG 111
QY 554 GGATGGCAGGTCCACCTGCAGACAGACCATGCCACAGGCG 596
Db 153 GGAGGCGCGTAGTAGTTCGATGTTGATCCGGTTGACATGCCG 111

RESULT 26
US-09-205-426-202/c
Sequence 202, Application US/09205426
Patent No. 6406704
GENERAL INFORMATION:
APPLICANT: Watson, James D.
APPLICANT: Tan, Paul L. J.
TITLE OF INVENTION: Compounds and Methods for Treatment and
TITLE OF INVENTION: Diagnosis of Mycobacterial Infections
FILE REFERENCE: 11000.1002c4
CURRENT APPLICATION NUMBER: US/09/205,426
CURRENT FILING DATE: 1998-12-04
EARLIER APPLICATION NUMBER: 09/095,855
EARLIER FILING DATE: 1998-06-11
EARLIER APPLICATION NUMBER: 08/997,362
EARLIER FILING DATE: 1997-12-23
EARLIER APPLICATION NUMBER: 08/873,970
EARLIER FILING DATE: 1997-06-12
EARLIER APPLICATION NUMBER: 08/705,347
EARLIER FILING DATE: 1996-08-29
NUMBER OF SEQ ID NOS: 208
SOFTWARE: FastSeq for Windows Version 3.0
SEQ ID NO 202
LENGTH: 570
TYPE: DNA
ORGANISM: Mycobacterium vaccae
US-09-205-426-202

Query Match 3.8%; Score 37.4; DB 4; Length 570;
Best Local Similarity 48.0%; Pred. No. 2;
Matches 107; Conservative 0; Mismatches 116; Indels 0; Gaps 0;
QY 374 CACCTCAGTCCCATGTGGACAGTCCAGGCTACAGATGACCTGGGAGCTGTGTCCTCAA 433
Db 333 CTCCTCGGTGAGCTGCGGGATCGACACCCGATGATGTTGCCGTGCTGCGATTGAC 274
QY 434 GCCAGTGTCTGAGCTGCGGGCACTTCACTCTCAGACAGAGAGGGGAGGCTGATCGGAA 553
Db 273 GCCAGTGTCTGAGCTGCGGGATCGACACCCGATGATGTTGCCGTGCTGCGATTGAC 274
QY 494 GGAGATGATCTGCCAGGCTCTCTCGGGCAGCCACCTATCACCAACAGCTGATCGGAA 553
Db 213 CTTGATCACCACCATGCGGCTCGGGCAGCTTGTGATCCGGTTGACATGCCG 111
QY 554 GGATGGCAGGTCCACCTGCAGACAGACCATGCCACAGGCG 596
Db 153 GGAGGCGCGTAGTAGTTCGATGTTGATCCGGTTGACATGCCG 111

RESULT 27
US-09-319-892-3
Sequence 3, Application US/09319892
Patent No. 6177616
GENERAL INFORMATION:
APPLICANT: BARTSCH, Klaus
APPLICANT: KRIETE, Guido
APPLICANT: BROER, Inge
APPLICANT: FUHLER, Alfred
TITLE OF INVENTION: NOVEL GENES CODING FOR AMINO ACID DEACETYLASES WITH
TITLE OF INVENTION: SPECIFICITY FOR N-ACETYL-L-PHOSPHINOTHRICIN, THEIR
FILE REFERENCE: 514412-2005
CURRENT APPLICATION NUMBER: US/09/319,892
CURRENT FILING DATE: 1999-06-14
EARLIER APPLICATION NUMBER: PCT/EP97/06755
EARLIER FILING DATE: 1997-12-03
EARLIER APPLICATION NUMBER: 19652284.6
EARLIER FILING DATE: 1996-12-16
NUMBER OF SEQ ID NOS: 4
SOFTWARE: PatentIn Ver. 2.0
SEQ ID NO 3
LENGTH: 1273
TYPE: DNA
ORGANISM: Stenotrophomonas maltophilia
US-09-319-892-3

Query Match 3.7%; Score 36.6; DB 3; Length 1273;
Best Local Similarity 45.0%; Pred. No. 4.3;
Matches 138; Conservative 0; Mismatches 169; Indels 0; Gaps 0;
QY 431 CAAGCCAGTGTCTGAGCTGCGGCCCACTTCACTCTGAGGACAGAGGGGAGGCCCCAG 490
Db 735 CGAGGCCACCAAGCTGGTCCCGCAGACGCTGAGCTGACAGGACCGGTGGCACCCTTAC 794
QY 491 GGTGAGATGATCTGCCAGGCGCTCTCTCGGGCAGCCACCTATCACCAACAGCTGATCGG 550
Db 795 GCTGGAGGTGCTGGACCTGATCGAGCGGCGATGAAGGCCCTGGCGGAGAGCATCTGCC 854
QY 551 GAAGGATGGCAGGCTCCACTGACGACAGACATGCGACAGGAGCTGCGCAACTTCTC 610
Db 855 GCGCATGACACGCGCTGCGAGTTGAGTTGCTGCGCAACTACCCGCCACCACCATCACTC 914
QY 611 CTTCTGCGAGCCAGACATCGGACTGTTCTGTTGCGCAGGCTGCAAAACACCACTATCT 670
Db 915 CGCCCGGAGGCGGATTCGACGCGCGCTCATGCGCCGAGGTGCTGGGCGAGGCCACGT 974
QY 671 CCAGCACAGCGCCCTCACAGTGGTGGCCCCAGGTGTGACCAAGAGATGAGGACTGGCA 730
Db 975 GCTGCCCCAGGAGCGCTCCATGGGCGCGGAGGACTTGGCTTTCATGCTGAAAAAGCC 1034

```
QY 731 GGGTCCC 737
Db 1035 CGGCGCC 1041

RESULT 28
US-09-727-238-3
; Sequence 3, Application US/09727238
; Patent No. 655733
; GENERAL INFORMATION:
; APPLICANT: Bartsch, et al.
; TITLE OF INVENTION: No. 655733el genes coding for amino acid deacetylases with speci
; FILE REFERENCE: acetyl-L-phosphinothricin, their isolation and their use.
; CURRENT APPLICATION NUMBER: 514412A-2005.1
; CURRENT FILING DATE: 2000-11-30
; PRIOR APPLICATION NUMBER: US/09/727,238
; PRIOR FILING DATE: 1999-06-14
; PRIOR APPLICATION NUMBER: PCT/EP97/06755
; PRIOR FILING DATE: 1997-12-03
; PRIOR APPLICATION NUMBER: 196 52 284.6
; PRIOR FILING DATE: 1996-12-06
; NUMBER OF SEQ ID NOS: 4
; SOFTWARE: PatentIn version 3.0
; SEQ ID NO 3
; LENGTH: 1273
; TYPE: DNA
; ORGANISM: Comamonas acidovorans
; NAME/KEY: CDS
; LOCATION: (1)...(1206)
; OTHER INFORMATION: coding sequence of the deacetylase gene from comamonas acidovorans
US-09-727-238-3
```

```
Query Match 3.7%; Score 36.6; DB 4; Length 1273;
Best Local Similarity 45.0%; Pred. No. 4.3;
Matches 138; Conservative 0; Mismatches 169; Indels 0; Gaps 0;

QY 431 CAAGCCAGTCTTGAGTGGCGGCCAATTCACCTGTCAGGACAGAGGGGCGGCCAG 490
Db 735 CGAGGGCCACAGCTGTGCTGCCACAGCGTGGAGCTGCAGGGCAGCGTGGCGACCTTCAC 794
QY 491 GGTGGAGATGATCTGCCAGCGCTCTCGGGCAGCCACCTATCACCAGCGCTGATCGG 550
Db 795 GCTGGAGGTGCTGGACCTGATCAGCGGGCGCATGAAAGGCCCTGGCGGAGAGCATCTGCGC 854
QY 551 GAAGGATGGGCGAGTCCACCTGTCAGCAGAGACCCATGCCAGCAGCGCTGCCAATCTTC 610
Db 855 GCGCATGACACGCGTGGAGTTCGAGTTCGTGCGCACTACCCGCCACCATCAACTC 914
QY 611 CTTCTGCGGAGCCAGCATCGGACTGGTGTGTCGCGAGGTGTCGAGGTCGCAACAGCCCAATGT 670
Db 915 CGCCCGGAGGCGGAGTTCGACGCGCGCTCATGGCGGAGGTGTCGGCGAGGCCAAGT 974
QY 671 CCAGCAGACGCGCTCACAGTGTGCGCCCGAGGTGTGACCAAGATGAGGACTGGCA 730
Db 975 GTGTCGCCAGAGGCGCTCCATGGGCGCCGAGGACTTCGCTTCTGCTGCAAGGCC 1034
QY 731 GGGTCCC 737
Db 1035 CGGCGCC 1041
```

```
RESULT 29
US-09-252-991A-11102
; Sequence 11102, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
```

```
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11102
; LENGTH: 801
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11102

Query Match 3.7%; Score 36.4; DB 4; Length 801;
Best Local Similarity 50.0%; Pred. No. 4.1;
Matches 91; Conservative 0; Mismatches 91; Indels 0; Gaps 0;

QY 601 CCAACTTCTCTTCTTCTGCGAGCCAGACATCGGACTGTTCTGTGTCAGGCTGCAACA 660
Db 279 CCAAGGCTTTGCGCGAGCGCTCGACAAGCTGTGCCGATCAGCGTCAAGGAAGCAGAGG 338
QY 661 ACGCCATGTCCAGCAGCAGCGCTCTACAGTGTGCCCCCAGGTGTTGACCAAGATGG 720
Db 339 ACGGCGACATCTCTGCTTCCGCTGCGCTGCTGCGCGCGCGGCGGCAAGCAGATGATGG 398
QY 721 AGGACTGGCAGGCTCCCTTGGAGAGCCCATCTTGCCTTTCGCGCTCTACAGGACACCC 780
Db 399 TCAGCGCGCGGCGCAGGTTGCGCATCTCTCCCGGCGAGAACGCTGAACTACAAGCCCT 458
QY 781 GC 782
Db 459 GC 460
```

```
RESULT 30
US-09-252-991A-11158
; Sequence 11158, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11158
; LENGTH: 1116
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11158
```

```
Query Match 3.7%; Score 36.4; DB 4; Length 1116;
Best Local Similarity 50.0%; Pred. No. 4.6;
Matches 91; Conservative 0; Mismatches 91; Indels 0; Gaps 0;

QY 601 CCAACTTCTCTTCTTCTGCGAGCCAGACATCGGACTGTTCTGTGTCAGGCTGCAACA 660
Db 683 CCAAGGCTTTGCGCGAGCGCTCGACAAGCTGTGCCGATCAGCGTCAAGGAAGCAGAGG 742
QY 661 ACGCCATGTCCAGCAGCAGCGCTCTACAGTGTGCCCCCAGGTGTTGACCAAGATGG 720
Db 743 ACGGCGACATCTCTGCTTCCGCTGCGCTGCTGCGCGCGCGGCGGCAAGCAGATGATGG 802
QY 721 AGGACTGGCAGGCTCCCTTGGAGAGCCCATCTTGCCTTTCGCGCTCTACAGGACACCC 780
Db 803 TCAGCGCGCGGCGCAGGTTGCGCATCTCTCCCGGCGAGAACGCTGAACTACAAGCCCT 862
QY 781 GC 782
Db 459 GC 460
```


Db 863 GC 864

RESULT 31

```

US-09-252-991A-11094/G
; Sequence 11094, Application US/09252991A
; Patent No. 6551795
; GENERAL INFORMATION:
; APPLICANT: Marc J. Rubenfield et al.
; TITLE OF INVENTION: NUCLEIC ACID AND AMINO ACID SEQUENCES RELATING TO PSEUDOMONAS
; TITLE OF INVENTION: AERUGINOSA FOR DIAGNOSTICS AND THERAPEUTICS
; FILE REFERENCE: 107196.136
; CURRENT APPLICATION NUMBER: US/09/252,991A
; CURRENT FILING DATE: 1999-02-18
; PRIOR APPLICATION NUMBER: US 60/074,788
; PRIOR FILING DATE: 1998-02-18
; PRIOR APPLICATION NUMBER: US 60/094,190
; PRIOR FILING DATE: 1998-07-27
; NUMBER OF SEQ ID NOS: 33142
; SEQ ID NO 11094
; LENGTH: 1584
; TYPE: DNA
; ORGANISM: Pseudomonas aeruginosa
US-09-252-991A-11094

```

	Query Match	3.7%;	Score 36.4;	DB 4;	Length 1584;
	Best Local Similarity	50.0%;	Pred. No. 5.2;		
	Matches	91;	Conservative	0;	Mismatches 91;
					Indels 0;
					Gaps 0;
QY	601	CCAACTTCTCCTTCCTGCCGAGCAGACATCGGACTGGTTCGTGTCGAGGCTCGAACA	660		
Db	839	CGAAGGCTTTGCGCGAGCGGCTCGACAAGGTGTCGGGATCAGCGTCAAGGAACGAGG	780		
QY	661	ACGCCAATGTCCTCAGCACAGCGCCCTCACAGTGTGCCCCAGGTGGTGACCCAGAGAATGG	720		
Db	779	ACGGCGACATCTTGGCGTCCGGGCTGGCGTGTGGGCGCGGCGGACGACGATGATGG	720		
QY	721	AGGACTGGCAGGTTCCCTCTGGAGGCCCAATCCCTTGGCTTTCGCCCTCTACGAGGACCC	780		
Db	719	TCGACGCGCCGCGCAGGTCGATCTCTCCCGCGACGAACGCGCTGAACTACAAAGCCCT	660		

RESULT 32

```

US-09-211-930-12
; Sequence 12, Application US/09211930
; Patent No. 5962265
; GENERAL INFORMATION:
; APPLICANT: Tyrell E. No. 5962265ris
; APPLICANT: William Craig Moore
; APPLICANT: David Shay Silberstein
; TITLE OF INVENTION: HUMAN SIGNAL TRANSDUCTION
; TITLE OF INVENTION: SERINE/THREONINE KINASE
; FILE REFERENCE: PHM. 70296
; CURRENT APPLICATION NUMBER: US/09/211,930
; CURRENT FILING DATE: 1998-12-15
; EARLIER APPLICATION NUMBER: GB 9726851.0
; EARLIER FILING DATE: 1997-12-19
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 12
; LENGTH: 2028
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-211-930-12

```

Query Match 3.7%; Score 36.4; DB 2; Length 2028;
Best Local Similarity 79.6%; Pred. No. 5.7;
Matches 43; Conservative 0; Mismatches 11; Indels 0; Gaps 0;

Qy	939	AAAAAGAGTGTGTTT	TAGCTCTCTT	GCCACAAAAA	AAAAAAAAAAAA	992
Db	1950	AAAAACAAGTGTGTTT	TAAAGAGCTCCCG	AAAAA	AAAAAAAAAAAA	200

RESULT 33

```

US-09-340-993-12
; Sequence 12, Application US/09340993
; Patent No. 6034228
; GENERAL INFORMATION:
; APPLICANT: Tyrell E. No. 6034228:ris
; APPLICANT: William Craig Moore
; APPLICANT: David Shay Silberstein
; TITLE OF INVENTION: HUMAN SIGNAL TRANSDUCTION SERINE/THREONINE KINASE
; FILE REFERENCE: PHM.70296.N1
; CURRENT APPLICATION NUMBER: US/09/340,993
; CURRENT FILING DATE: 1999-06-25
; EARLIER APPLICATION NUMBER: GB 9726851.0 & US 09/211,930
; EARLIER FILING DATE: 1997-12-19 & 1998-12-15
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 12
; LENGTH: 2028
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-340-993-12

```

Query Match	3.7%	Score 36.4;	DB 3;	Length 2028;
Best Local Similarity	79.8%;	Pred. No. 5.7;		
Matches 43;	Conservative	0;	Mismatches 11;	Indels 0;
Gaps	0;			
QY	939	AAATGAGTGTGTTT	TAGTGCTCTT	CGCCAAAAA
		AAAAAAAAAAAAAAAA	AAAAAAAAAAAA	992
Db	1950	AAACACAAGTGT	TTTTAAGAAGCT	CCCGAAAAA
		AAAAAAAAAAAAAAAA	AAAAAAAAAAAA	2003

RESULT 34

```

US-09-468-442-12
; Sequence 12, Application US/09458442
; Patent No. 6300098
; GENERAL INFORMATION:
; APPLICANT: Tyrell E. No. 6300098ris
; APPLICANT: William Craig Moore
; APPLICANT: David Shay Silberstein
; TITLE OF INVENTION: HUMAN SIGNAL TRANSDUCTION SERINE/THREONINE KINASE
; FILE REFERENCE: PHM.70296.N1
; CURRENT APPLICATION NUMBER: US/09/468,442
; CURRENT FILING DATE: 1999-12-21
; EARLIER APPLICATION NUMBER: US 09/340,993
; EARLIER FILING DATE: 1999-06-25
; EARLIER APPLICATION NUMBER: GB 9726851.0
; EARLIER FILING DATE: 1997-12-19
; EARLIER APPLICATION NUMBER: US 09/211,930
; EARLIER FILING DATE: 1998-12-15
; NUMBER OF SEQ ID NOS: 14
; SOFTWARE: FastSeq for Windows Version 3.0
; SEQ ID NO 12
; LENGTH: 2028
; TYPE: DNA
; ORGANISM: Mus musculus
US-09-468-442-12

```

	Query Match	3.7%	Score 36.4;	DB 4;	Length 2028;
	Best Local Similarity	79.6%	Pred. No. 5.7;		
	Matches 43;	Conservative 0;	Mismatches 11;	Indels 0;	Gaps 0;
QY	939	AAAATGAGTGTGTTTTAGCTGCTCTTCCACAAAAA	AAAAAAAAAAAAAAAAAAAA	992	
DB	1950	AAAAACAAGTGTGTTTTAAGTAGAGTCCCGAAAAA	AAAAAAAAAAAAAAAAAAAA	2003	

RESULT 35

US-09-252-991A-5589/c


```

1 STREET: 7 Skyline Drive
2 CITY: Hawthorne
3 STATE: NY
4 COUNTRY: USA
5 ZIP: 10532
6
7 COMPUTER READABLE FORM:
8 MEDIUM TYPE: Floppy disk
9 COMPUTER: IBM PC compatible
10 OPERATING SYSTEM: PC-DOS/MS-DOS
11 SOFTWARE: PatentIn Release #1.0, Version #1.25
12 CURRENT APPLICATION DATA:
13 APPLICATION NUMBER: US/08/457,342
14 FILING DATE: 01-JUN-1995
15 CLASSIFICATION: 424
16 PRIOR APPLICATION DATA:
17 APPLICATION NUMBER: US 08/457,205
18 FILING DATE: 01-JUN-1995
19 APPLICATION NUMBER: 08/258,261
20 FILING DATE: 08-Jun-1994
21 ATTORNEY/AGENT INFORMATION:
22 NAME: Elmer, James Scott
23 REGISTRATION NUMBER: 36,129
24 REFERENCE/DOCKET NUMBER: CGC 1506/CIP3
25 TELECOMMUNICATION INFORMATION:
26 TELEPHONE: 919-541-8614
27 TELEFAX: 919-541-8689
28 INFORMATION FOR SEQ ID NO: 6:
29 SEQUENCE CHARACTERISTICS:
30 LENGTH: 28958 base pairs
31 TYPE: nucleic acid
32 STRANDEDNESS: single
33 TOPOLOGY: linear
34 MOLECULE TYPE: DNA (genomic)
35 HYPOTHETICAL: NO
36 ANTI-SENSE: NO
37 US-08-457-342-6

```

Query Match	3.6%	Score 35.8;	DB 1;	Length 28958;
Best Local Similarity	52.3%;	Pred. No. 21;		
Matches	79;	Conservative 0;	Mismatches 72;	Indels 0; Gaps 0;
QY	56	CCGAGGCGATGGGCTCCCTGGCGTGTTCTCTCTGGCGCGTCTGGCTCCAGAGCTTCTC	115	
Db	4498	CAGGGCCATTGGCGCGGTGAGGCTTGGCTCGCGCGCTCTGGTTACGCGCGAGCGCTC	4439	
QY	116	CRAAGCACGGAGGAGAAATTAACCTCTGTGGTCTCCATTGCCTACAAAGTCTCTGGAAGT	175	
Db	4438	GAAGGACGGCGAGGACGGGATGACCGCTTTTGGACGGCATCGAGAGCGCTTCGAGCAGGA	4379	
QY	176	TTTTCGCGAAGCGCGTGGGTGCTCATACCC	206	
Db	4378	GCATCCGGCGCGCTCTGGCCACCGCGAACC	4348	

```

RESULT 44
US-08-457-646A-6/c
; Sequence 6, Application US/08457646A
; Patent No. 5679560
;
; GENERAL INFORMATION:
;
; APPLICANT: Schupp, Thomas
; APPLICANT: Ligon, James M.
; APPLICANT: Beck, James Joseph
; APPLICANT: Hill, Dwight Steven
; APPLICANT: Ryals, John Andrew
; APPLICANT: Gaffney, Thomas Deane
; APPLICANT: Lam, Stephen Ting
; APPLICANT: Hammer, Phillip E.
; APPLICANT: Uknes, Scott Joseph
;
; TITLE OF INVENTION: Genes for the synthesis of
; TITLE OF INVENTION: antipathogenic substances
; NUMBER OF SEQUENCES: 22
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ciba-Geigy Corporation
;

```

```

1 STREET: 7 Skyline Drive
2 CITY: Hawthorne
3 STATE: NY
4 COUNTRY: USA
5 ZIP: 10532
6
7 COMPUTER READABLE FORM:
8 MEDIUM TYPE: Floppy disk
9 COMPUTER: IBM PC compatible
10 OPERATING SYSTEM: PC-DOS/MS-DOS
11 SOFTWARE: PatentIn Release #1.0, Version #1.25
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13 CURRENT APPLICATION DATA:
14 APPLICATION NUMBER: US/03/457,646A
15 FILING DATE: 01-JUN-1995
16 CLASSIFICATION: 530
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18 PRIOR APPLICATION DATA:
19 APPLICATION NUMBER: US 08/457,205
20 FILING DATE: 01-JUN-1995
21 APPLICATION NUMBER: 08/258,261
22 FILING DATE: 08-Jun-1994
23 ATTORNEY/AGENT INFORMATION:
24 NAME: Elmer, James Scott
25 REGISTRATION NUMBER: 36,129
26 REFERENCE/DOCKET NUMBER: CGC 1506/CIP3
27 TELECOMMUNICATION INFORMATION:
28 TELEPHONE: 919-541-8614
29 TELEFAX: 919-541-8689
30 INFORMATION FOR SEQ ID NO: 6:
31
32 SEQUENCE CHARACTERISTICS:
33 LENGTH: 28958 base pairs
34 TYPE: nucleic acid
35 STRANDEDNESS: single
36 TOPOLOGY: linear
37 MOLECULE TYPE: DNA (genomic)
38 HYPOTHETICAL: NO
39 ANTI-SENSE: NO
40 US-08-457-646A-6
41
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43 Best Local similarity 52.3%; Pred. No. 21;
44 Matches 79; Conservative 0; Mismatches 72;
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47 DB 4498 CAGGGCCATTGGGGCCGGGTGAGCGCTTGGCTCCGGCCGTGC
48 QY 116 CAAGGACCGGAGGAGAAATATCCCTGTGGTCTCCATTG
49 DB 4438 GAAGGACGGCGAGGACGGGATACCGTTTGGACGGCATCG
50 QY 176 TTTTCCCCAAAGGCCGCTGGGTGCTATAAACC 206
51 DB 4378 GCATCCCGCGCGCCCTCGGCGCCGAGCCGAACC 4348

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1  RESULT 45
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3  Sequence 6, Application US/08458076A
4  Patent No. 5698425
5  GENERAL INFORMATION:
6  APPLICANT: Schupp, Thomas
7  APPLICANT: Ligon, James M.
8  APPLICANT: Beck, James Joseph
9  APPLICANT: Hill, Dwight Steven
10 APPLICANT: Ryalls, John Andrew
11 APPLICANT: Gaffney, Thomas Deane
12 APPLICANT: Lam, Stephen Ting
13 APPLICANT: Hammer, Phillip E.
14 APPLICANT: Uknes, Scott Joseph
15 TITLE OF INVENTION: Genes for the synthesis of
16 TITLE OF INVENTION: antipathogenic substances
17 NUMBER OF SEQUENCES: 22
18 CORRESPONDENCE ADDRESS:
19 ADDRESSER: Ciba-Geigy Corporation
20

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STREET: 7 Skyline Drive
CITY: Hawthorne
STATE: NY
COUNTRY: USA
ZIP: 10532
COMPUTER READABLE FORM:
MEDIUM TYPE: Floppy disk
COMPUTER: IBM PC compatible
OPERATING SYSTEM: PC-DOS/MS-DOS
SOFTWARE: Patent In Release #1.0, Version #1.25
CURRENT APPLICATION DATA:
APPLICATION NUMBER: US/08/458,076A
FILING DATE: 01-JUN-1995
CLASSIFICATION: 435
PRIOR APPLICATION DATA:
APPLICATION NUMBER: US 08/457,205
FILING DATE: 01-JUN-1995
APPLICATION NUMBER: 08/258,261
FILING DATE: 08-JUN-1994
ATTORNEY/AGENT INFORMATION:
NAME: Elmer, James Scott
REGISTRATION NUMBER: 36,129
REFERENCE/DOCKET NUMBER: CGC 1506/CIP3
TELECOMMUNICATION INFORMATION:
TELEPHONE: 919-541-8614
TELEFAX: 919-541-8689
INFORMATION FOR SEQ ID NO: 6:
SEQUENCE CHARACTERISTICS:
LENGTH: 28958 base pairs
TYPE: nucleic acid
STRANDEDNESS: single
TOPOLOGY: linear
MOLECULE TYPE: DNA (genomic)
HYPOTHETICAL: NO
ANTI-SENSE: NO
US-08-458-076A-6

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Query Match          3.6%; Score 35.8; DB 1; Length 28958;
Best Local Similarity 52.3%; Pred. No. 21;
Matches 79; Conservative 0; Mismatches 72; Indels 0;

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4498	CAGGGCCATTGGCGCGGTGAGCCCTTGGCTCCGGCGCTCCTGGTTGACGCGGAGCCTC	4439
Db		
116	CAAGGCACGGAGGAGAAATTACCCCTGTGGTCTCCATTGCCATCAAAATCCTCGAAGT	175
QY		
4438	GAAGGACGCGAGGACGGATGACCGTTTGGACGGCATCGAGAGCGCTCGAGCAGGA	4379
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QY		
4378	GCATCCGCGCGCTCGGCCAGCCGAACT	4348
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RESULT 46
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; Sequence 4, Application US/08764233A
; Patent No. 5716849
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; GENERAL INFORMATION:
;
; APPLICANT: Ligon, James M.
; APPLICANT: Schupp, Thomas
; APPLICANT: Beck, James J.
; APPLICANT: Hill, Dwight S.
; APPLICANT: Neff, Snezana
; APPLICANT: Ryals, John A.
;
; TITLE OF INVENTION: Genes For The Biosynthesis Of Soraphen
;
; NUMBER OF SEQUENCES: 10
;
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ciba-Geigy Corporation
; STREET: 520 White Plains Road, P.O. Box 2005
; CITY: Tarrytown
; STATE: NY
; COUNTRY: USA

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ZIP: 10591
 COMPUTER READABLE FORM:
 MEDIUM TYPE: Floppy disk
 COMPUTER: IBM PC compatible
 OPERATING SYSTEM: PC-DOS/MS-DOS
 SOFTWARE: PatentIn Release #1.0, Version #1.30
 CURRENT APPLICATION DATA:
 APPLICATION NUMBER: US/08/764,233A
 FILING DATE:
 CLASSIFICATION: 435
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/729,214
 FILING DATE: 09-OCT-1996
 PRIOR APPLICATION DATA:
 APPLICATION NUMBER: US 08/258,261
 FILING DATE: 08-JUN-1994
 ATTORNEY/AGENT INFORMATION:
 NAME: Meigs, J. Timothy
 REGISTRATION NUMBER: 38,241
 REFERENCE/DOCKET NUMBER: 1506/CIP6
 TELECOMMUNICATION INFORMATION:
 TELEPHONE: (919) 541-8587
 TELEFAX: (919) 541-8689
 INFORMATION FOR SEQ ID NO: 4:
 SEQUENCE CHARACTERISTICS:
 LENGTH: 28958 base pairs
 TYPE: nucleic acid
 STRANDEDNESS: single
 TOPOLOGY: linear
 MOLECULE TYPE: DNA (genomic)
 ORIGINAL SOURCE:
 ORGANISM: Sorangium cellulosum
 IMMEDIATE SOURCE:
 CLONE: p98/1
 US-08-764-233A-4

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QY	176	TTTCCCAGAGCGCGTGGTGTCTATAACC	206		
Db	4378	GCATCCGCGCGCGCTCGGCCACGCCGAAAC	4348		

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Qy	116	CAAGGCACGGAGGAGAAATTACCCCTGTGGTCTCCATTGCCACAAAGTCTCTGGAAGT	175
Db	4438	GAAGGACGGCGAGGACGGGTATGACCTTTTGGACGGCATCGGAGGCGGCTCGAGCAGGA	4379
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RESULT 47
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? Sequence 6, Application US/08457335A
? Patent No. 5723759
? GENERAL INFORMATION:
? APPLICANT: Schnupp, Thomas
? APPLICANT: Ligon, James M.
? APPLICANT: Beck, James Joseph
? APPLICANT: Hill, Dwight Steven
? APPLICANT: Ryals, John Andrew
? APPLICANT: Gaffney, Thomas Deane
? APPLICANT: Lam, Stephen ting
? APPLICANT: Hammer, Phillip E.
? APPLICANT: Uknes, Scott Joseph
? TITLE OF INVENTION: Genes for the synthesis of
? TITLE OF INVENTION: antipathogenic substances
? NUMBER OF SEQUENCES: 22
? CORRESPONDENCE ADDRESS:
? ADDRESSEE: Ciba-Geigy Corporation
? STREET: 7 Skyline Drive

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; CITY: Hawthorne
; STATE: NY
; COUNTRY: USA
; ZIP: 10532
;
; COMPUTER READABLE FORM:
; MEDIUM TYPE: Floppy disk
; COMPUTER: IBM PC compatible
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; APPLICATION NUMBER: US/08/457,335A
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; PRIOR APPLICATION DATA:
; APPLICATION NUMBER: US 08/457,205
; FILING DATE: 01-JUN-1995
; APPLICATION NUMBER: 08/258,261
; FILING DATE: 08-JUN-1994
; ATTORNEY/AGENT INFORMATION:
; NAME: Elmer, James Scott
; REGISTRATION NUMBER: 36,129
; REFERENCE/DOCKET NUMBER: CGC 1506/CIP3
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 919-541-8614
; TELEFAX: 919-541-8689
; INFORMATION FOR SEQ ID NO: 6:
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; LENGTH: 28958 base pairs
; TYPE: nucleic acid
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; HYPOTHETICAL: NO
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; Db 4438 GAAGGACGGCGGAGGACGGGATGACCGTTTGGACGGCATCGGAGGCGGCTCGAGCAGGA 4379
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; QY 176 TTTCCTCCAAAGCGCGCTGGGTGCTCATTAACC 206
; Db 4378 GCATCCCGCGCCCTCGGCCCGAGCCCGAACC 4348
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; RESULT 48
; US-08-729-214-6/c
; Sequence 6, Application US/08/29214
; Patent No. 5817502
; GENERAL INFORMATION:
; APPLICANT: Ligon, James M.
; APPLICANT: Hill, Dwight Steven
; APPLICANT: Ryals, John Andrew
; APPLICANT: Hammer, Phillip E.
; APPLICANT: van Pee, Karl-Heinz
; APPLICANT: Kirner, Sabine
; TITLE OF INVENTION: Genes for the synthesis of
; TITLE OF INVENTION: antipathogenic substances
; NUMBER OF SEQUENCES: 27
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: Ciba-Geigy Corporation
; STREET: 520 White Plains Road
; CITY: Tarrytown
; STATE: NY
; COUNTRY: USA
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; ZIP: 10591
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; CURRENT APPLICATION DATA:
; APPLICATION NUMBER: US/08/729,214
; FILING DATE: TBA
; CLASSIFICATION: 435
; ATTORNEY/AGENT INFORMATION:
; NAME: Meigs, J. Timothy
; REGISTRATION NUMBER: 38,241
; REFERENCE/DOCKET NUMBER: CGC 1506/CIP5
; TELECOMMUNICATION INFORMATION:
; TELEPHONE: 919-541-8587
; TELEFAX: 919-541-8689
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; Matches 79; Conservative 0; Mismatches 72; Indels 0; Gaps 0;
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; Db 4438 GAAGGACGGCGGAGGACGGGATGACCGTTTGGACGGCATCGGAGGCGGCTCGAGCAGGA 4379
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; QY 176 TTTCCTCCAAAGCGCGCTGGGTGCTCATTAACC 206
; Db 4378 GCATCCCGCGCCCTCGGCCCGAGCCCGAACC 4348
;
; RESULT 49
; US-09-028-934-6/c
; Sequence 6, Application US/09028934
; Patent No. 6117670
; GENERAL INFORMATION:
; APPLICANT: Ligon, James M.
; APPLICANT: Hill, Dwight S.
; APPLICANT: Lam, Steven T.
; APPLICANT: Hammer, Phillip E.
; APPLICANT: van Pee, Karl-Heinz
; APPLICANT: Kirner, Sabine
; APPLICANT: Young, Thomas R.
; TITLE OF INVENTION: Pyrolytic Biosynthesis Genes and Uses
; TITLE OF INVENTION: Thereof
; NUMBER OF SEQUENCES: 37
; CORRESPONDENCE ADDRESS:
; ADDRESSEE: No. 6117670artis Corporation
; STREET: 3054 Cornwallis Road
; CITY: Research Triangle Park
; STATE: NC
; COUNTRY: USA
; ZIP: 27709
; COMPUTER READABLE FORM:
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; COMPUTER: IBM PC compatible
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; SOFTWARE: Patentin Release #1.0, Version #1.30
; CURRENT APPLICATION DATA:
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GenCore version 5.1.6
Copyright (c) 1993 - 2004 CompuGen Ltd.

OM nucleic - nucleic search, using sw model

Run on: February 9, 2004, 12:59:37 ; Search time 328 Seconds

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Title: US-09-990-726-222

Perfect score: 992

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Searched: 2552756 seqs, 1349719017 residues

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Maximum Match 100%

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Pred. No. is the number of results predicted by chance to have a score greater than or equal to the score of the result being printed, and is derived by analysis of the total score distribution.

SUMMARIES

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6	992	100.0	992	25	ABX81154 Novel human secret
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8	992	100.0	992	25	ABX77855 Human PRO polynucl

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22	149.6	15.1	150	19	AAZ12009 Human NOV93a cDNA.
23	47.2	4.8	1272	24	ABX97141 Human G-protein co
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25	47	4.7	87350	18	AAZ83003 DNA encoding novel
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29	43	4.3	509	24	ABT07672 Genomic sequence e
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C 89	36.2	3.6	295	22	AAS29052	cDNA encoding for
C 90	36.2	3.6	295	24	ABS68192	cDNA encoding huma
C 91	36.2	3.6	304	22	AAS29128	cDNA encoding huma
C 92	36.2	3.6	304	24	ABS68268	cDNA encoding huma
C 93	36.2	3.6	359	22	AAK23694	Human breast cance
C 94	36.2	3.6	590	24	ABL59497	EST related to hum
C 95	36.2	3.6	1641	20	AAZ10823	Choline oxidase (C
C 96	36.2	3.6	2821	22	ABL66075	Triticum aestivum
C 97	36.2	3.6	2920	20	ABL35012	Rat cDNA isolated
C 98	36.2	3.6	541	20	AAK87415	Hepatocellular car
C 99	36.2	3.6	839	22	AAK86484	Maize Mac20 coding
C 100	36.2	3.6	1127	22	AAK07899	Human secreted pro
C 101	36.2	3.6	1128	22	AAK07835	Human secreted pro
C 102	36.2	3.6	1805	21	AAZ58260	Soybean histone de
C 103	36.2	3.6	1975	24	ABL90817	Human polynucleoti
C 104	36.2	3.6	2101	21	AAK93330	Human secreted pro
C 105	36.2	3.6	2101	22	AAF32713	Human secreted pro
C 106	36.2	3.6	2101	25	ABZ73609	Secreted protein-e
C 107	36.2	3.6	2101	25	ABZ67203	Human secreted pro
C 108	36.2	3.6	2625	21	AAZ31922	Human mahogany pro
C 109	36.2	3.6	2626	16	AAQ94582	Modified Tag DNA-p
C 110	36.2	3.6	10579	22	ABA20014	Human nervous syst
C 111	36.2	3.6	10579	21	AAK70045	Human immune/haema
C 112	36.2	3.6	21185	21	AAK63350	Streptomyces globi
C 113	36.2	3.6	28995	22	ABZ20015	Human nervous syst
C 114	36.2	3.6	28995	22	AAK70046	Human immune/haema
C 115	36.2	3.6	28995	22	AAK79967	Human immune/haema
C 116	36.2	3.6	28995	22	AAK85213	Human immune/haema
C 117	36.2	3.6	63164	21	AAK63348	Streptomyces globi
C 118	36.2	3.6	4403765	22	AAI99683	Mycobacterium tube
C 119	36.2	3.6	4411529	22	AAI99682	Mycobacterium tube
C 120	35.8	3.6	879	24	ABN98766	Arabidopsis thalia
C 121	35.8	3.6	1132	21	AAK76120	Human ORPX ORF1675
C 122	35.8	3.6	1725	24	ABA95144	Human lysyl-oxidase
C 123	35.8	3.6	2328	24	AAI67786	Human lysyl oxidase
C 124	35.8	3.6	28598	17	AAK06769	Sorangium cellulos
C 125	35.8	3.6	28598	18	AAK89956	Sorangium cellulos
C 126	35.8	3.6	28958	21	AAK75299	DNA sequence of So
C 127	35.8	3.6	43377	19	AAV05287	The soraphen biosy
C 128	35.8	3.6	144460	21	AAZ93815	Olfactory receptor
C 129	35.8	3.6	1503841	24	ABT00010	Human neuregulin 1
C 130	35.8	3.6	1503841	24	ABT01503	Human neuregulin 1
C 131	35.8	3.6	1503900	22	AAK95240	Human neuregulin-1
C 132	35.8	3.6	1503900	22	AAK96733	Human neuregulin-1
C 133	35.6	3.6	366	21	AAZ92717	Human wild-type cy
C 134	35.6	3.6	387	21	AAZ92716	Human wild-type cy
C 135	35.6	3.6	500	21	AAK54277	Arabidopsis thalia
C 136	35.6	3.6	577	24	ABL66282	Lung cancer relate
C 137	35.6	3.6	577	24	ABL67527	Thyroid cancer rel
C 138	35.6	3.6	577	24	ABL67787	Oesophagus cancer
C 139	35.6	3.6	581	21	AAZ35658	Human cystatin E
C 140	35.6	3.6	588	21	AAZ60085	Human cystatin E
C 141	35.6	3.6	588	22	AAD20097	Human cystatin E
C 142	35.6	3.6	588	24	AAK43452	Human gene express
C 143	35.6	3.6	591	24	AAZ35213	Human gene express
C 144	35.6	3.6	598	18	AAK06068	Human cysteine pro
C 145	35.6	3.6	1536	23	AAK51510	Pseudomonas aerugi
C 146	35.6	3.6	2046	24	ABQ54507	Human ovarian anti
C 147	35.6	3.6	16759	24	ABL36297	Human lysosomal ac
C 148	35.4	3.6	294	23	ABV61758	Human prostate exp
C 149	35.4	3.6	356	23	ABV10788	Human prostate exp
C 150	35.4	3.6	361	23	ABV01619	Human prostate exp

ALIGNMENTS

RESULT 1

AAK58373
 ID AAC58373 standard; cDNA; 992 BP.
 XX
 AC AAC58373;
 XX
 DT 29-JAN-2001 (first entry)
 XX
 DE Human PRO809 nucleotide sequence SEQ ID NO:22.

Human; tumour; diagnosis; neoplastic disease; neoplastic cell growth;
 proliferation; tumorigenesis; identification; cancer; cytostatic;
 neoplastic; neuroprotective; antiinflammatory; immunosuppressive;
 immunostimulant; antiangiogenic; leukaemia; lymphoid malignancy;
 neuronal disorder; glial disorder; astrocytal disorder; angiogenic;
 hypothalamic disorder; glandular disorder; macrophagal disorder;
 epithelial disorder; stromal disorder; blastocoelec disorder;
 inflammatory disorder; immunologic disorder; ss.

OS Homo sapiens.

XX WO2000053755-A2.

XX 14-SEP-2000.

XX 06-JAN-2000; 2000WO-US00376.

XX 08-MAR-1999; 99WO-US05028.

XX 02-JUN-1999; 99WO-US12252.

XX 23-JUN-1999; 99US-0141037.

XX 07-JUL-1999; 99US-0143048.

XX 26-JUL-1999; 99US-0145698.

XX 30-NOV-1999; 99WO-US28313.

XX 20-DEC-1999; 99WO-US30911.

XX 05-JAN-2000; 2000WO-US00219.

(GETH) GENENTECH INC.

XX Ashkenazi AJ, Baker KP, Goddard A, Gurney AL, Hillan KJ, Roy MA;
 XX Watanabe CK, Wood WI;
 XX WPI; 2000-572270/53.

XX P-PSDB; AAB24063.

XX Thirty PRO polynucleotides encoding PRO polypeptides, useful in the
 XX treatment, diagnosis and prevention of cancer -
 XX Claim 50; Fig 13; 286pp; English.

XX The present invention describes an isolated antibody that binds to

one of the human PRO proteins designated PRO212, PRO290, PRO341, PRO535,
 PRO619, PRO717, PRO809, PRO830, PRO848, PRO943, PRO1005, PRO1009,
 PRO1025, PRO1030, PRO1097, PRO1107, PRO1111, PRO1153, PRO1182, PRO1184,
 PRO1187, PRO1281, PRO23, PRO39, PRO834, PRO1317, PRO1710, PRO2094,
 PRO2145 OR PRO2198. PRO antagonists can be used to inhibit tumour cell
 growth. The PRO polypeptides and nucleotides are useful in the
 treatment, diagnosis and prevention of cancer. The antibodies and other
 anti-tumour compounds maybe used to treat various conditions, including
 those characterised by overexpression and/or activation of the amplified
 PRO genes. Exemplary conditions or disorders to be treated with such
 antibodies and other compounds include benign or malignant tumours
 (e.g., renal, liver, kidney, bladder, breast, gastric, thyroid, ovarian,
 colorectal, prostate, pancreatic, lung, vulva, thyroid, hepatic
 carcinomas, sarcomas, glioblastomas, and various disorders such as neuronal,
 leukaemias and lymphoid malignancies, other disorders such as neuronal,
 glial, astrocytal, hypothalamic and other glandular, macrophagal,
 epithelial, stromal and blastocoelec disorders, and inflammatory,
 angiogenic and immunologic disorders. AAC58242 to AAC58366 represent PCR
 primers and hybridisation probes used in the isolation of the human PRO
 sequences. AAC58367 to AAC58396 and AAB24057 to AAB24089 represent human
 PRO polynucleotide and protein sequences given in the exemplification of
 the present invention.

XX	Sequence	992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;
Query Match	100.0%;	Score 992; DB 21; Length 992;
Best Local Similarity	100.0%;	Pred. No. 2.7e-236;
Matches	992; Conservative 0;	Mismatches 0; Indels 0; Gaps 0;
Qy	1	GGCAGAGCCAGGAACCTAGGAGGTTCTCAGTCGCCGAGCAGAGGCGCTTACACCCACCCAG 60
Db	1	GGCAGGAGCCAGGAACCTAGGAGGTTCTCAGTCGCCGAGCAGAGGCGCTTACACCCACCCAG 60
Qy	61	GCATGGGGCTCCCTGGGCTGTTCTGCTTGGCGGTGCTGGCTGCAGCAGGCTTCTTCCAGG 120
Db	61	GCATGGGGCTCCCTGGGCTGTTCTGCTTGGCGGTGCTGGCTGCAGCAGGCTTCTTCCAGG 120
Qy	121	CACGGGAGGAAGAAATTACCCCTGTGGTCTCCATTGGCTCAAAAGTCTCTGGAAGTTTTC 180
Db	121	CACGGGAGGAAGAAATTACCCCTGTGGTCTCCATTGGCTCAAAAGTCTCTGGAAGTTTTC 180
Qy	181	CCAAAGCCGCTGGGTGCTCATACCTGCTGTGCACCCAGCCACCCAGCCACCGCCATCACCT 240
Db	181	CCAAAGCCGCTGGGTGCTCATACCTGCTGTGCACCCAGCCACCCAGCCACCGCCATCACCT 240
Qy	241	ATTCCCTCTGTGGAAACCAAGAACATCAAGTGTGGCAAGAGGTGGTGAAGACCCACGAGC 300
Db	241	ATTCCCTCTGTGGAAACCAAGAACATCAAGTGTGGCAAGAGGTGGTGAAGACCCACGAGC 300
Qy	301	CGGCTCTCTTCAACCTCAAGCTCACACTCAAGTCCAGTCCAGACCTGCTCACCTACTTCT 360
Db	301	CGGCTCTCTTCAACCTCAAGCTCAAGTCCAGTCCAGACCTGCTCACCTACTTCT 360
Qy	361	GCGGGGGCTCTCCACCTCAGGTCGCCATGTGACAGTGCAGGCTACAGATGCACTGGG 420
Db	361	GCGGGGGCTCTCCACCTCAGGTCGCCATGTGACAGTGCAGGCTACAGATGCACTGGG 420
Qy	421	AGCTGTGGTTCAGCCAGTGTCTGAGCTGCGGGCCAACTTCACTCTGCAGGACAGAGGGG 480
Db	421	AGCTGTGGTTCAGCCAGTGTCTGAGCTGCGGGCCAACTTCACTCTGCAGGACAGAGGGG 480
Qy	481	CAGSCCCAGGGTGGAGATGATCTGCCAGAGGCTCCTGGCGAGCGCCACCTTATCACCAACA 540
Db	481	CAGSCCCAGGGTGGAGATGATCTGCCAGAGGCTCCTGGCGAGCGCCACCTTATCACCAACA 540
Qy	541	GCTGTATCGGGAAGGATGGGAGGTCCACCTGCAGCAGAGACCATGCCACAGCAGGCGTG 600
Db	541	GCTGTATCGGGAAGGATGGGAGGTCCACCTGCAGCAGAGACCATGCCACAGCAGGCGTG 600
Qy	601	CGAACTTCTCCTCTGTCGAGGACAGACATCGGACTGGTCTGGTGCAGGCTGCAAAACA 660
Db	601	CGAACTTCTCCTCTGTCGAGGACAGACATCGGACTGGTCTGGTGCAGGCTGCAAAACA 660
Qy	661	ACGCCAATGTCAGCAGACAGCGCTCAGTGTGCCCCAGTGTGTGACAGAGAGATGG 720
Db	661	ACGCCAATGTCAGCAGACAGCGCTCAGTGTGCCCCAGTGTGTGACAGAGAGATGG 720
Qy	721	AGGACTGGCAGGGTCCCTGGAGAGCCCCATCTTGGCTTTGCCGCTCTACAGAGAGCACCC 780
Db	721	AGGACTGGCAGGGTCCCTGGAGAGCCCCATCTTGGCTTTGCCGCTCTACAGAGAGCACCC 780
Qy	781	GCGCTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGGATAGGGAATGGGGAGGTTCAGAGGAC 840
Db	781	GCGCTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGGATAGGGAATGGGGAGGTTCAGAGGAC 840
Qy	841	GCAAGCAGCAGCCATGTAGATGAACCGTTCAGAGAGCCAGCAAGCAGAGAGGACTGCA 900
Db	841	GCAAGCAGCAGCCATGTAGATGAACCGTTCAGAGAGCCAGCAAGCAGAGAGGACTGCA 900
Qy	901	GGCCATCAGCGTGCACGTGTTCTGATTTGGAGTTCATGCAAAATCAGTGCTTTTAGTGC 960
Db	901	GGCCATCAGCGTGCACGTGTTCTGATTTGGAGTTCATGCAAAATCAGTGCTTTTAGTGC 960
Qy	961	TCTTGGCACAAAAAATAAAAAAAAAAAAAA 992

CC and DNA. They may also be used to produce transgenic animals which are
 CC used to develop and screen therapeutically useful reagents. The PRO
 CC nucleotide and protein sequence can be used for tissue typing and in
 CC treating cancer. Anti-PRO antibodies can be used in diagnostic assays.
 CC AAF44270 to AAF44470 represent PCR primers and hybridisation probes used
 CC in the isolation of human PRO sequences. AAF44087 to AAF44269 and
 CC AAB65154 to AAB65300 represent human PRO polynucleotide and protein
 CC sequences given in the exemplification of the present invention.
 XX
 SQ Sequence 992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;

Query Match 100.0%; Score 992; DB 22; Length 992;
 Best Local Similarity 100.0%; Pred. No. 2.7e-236;
 Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCACGAGCCAGGAACCTAGGAGTTCTCACTGCCGAGCAGAGGCCCTACACCCACCGAG 60
 DB 1 GGCACGAGCCAGGAACCTAGGAGTTCTCACTGCCGAGCAGAGGCCCTACACCCACCGAG 60

QY 61 GCATGGGGTCCCTGGCTGTTCTGCTGGCGGTGCTGCTGCCAGCAGCTTCTCCAGG 120
 DB 61 GCATGGGGTCCCTGGCTGTTCTGCTGGCGGTGCTGCTGCCAGCAGCTTCTCCAGG 120

QY 121 CACGGGAGGAATAATACCCCTGTGTCCTCATTCCTCAAAAGTCTCGAAGTTTTC 180
 DB 121 CACGGGAGGAATAATACCCCTGTGTCCTCATTCCTCAAAAGTCTCGAAGTTTTC 180

QY 181 CCAAGGGCGGTGGTGTCTATACCTGTGTGACCCCGCAGCAGCCGCGCCATCACT 240
 DB 181 CCAAGGGCGGTGGTGTCTATACCTGTGTGACCCCGCAGCAGCCGCGCCATCACT 240

QY 241 ATTCCCTCTGTGGACCAAGAACATCAAGGTGGCCAGAGTGTGTGAAGCCCAAGC 300
 DB 241 ATTCCCTCTGTGGACCAAGAACATCAAGGTGGCCAGAGTGTGTGAAGCCCAAGC 300

QY 301 CGGCTCTCTCAACCTCAACCTCACTCAAGTCCAGCTCCAGCTGCTCACTACTTCT 360
 DB 301 CGGCTCTCTCAACCTCAACCTCACTCAAGTCCAGCTCCAGCTGCTCACTACTTCT 360

QY 361 GCCGGGCTCTCACTCAAGTGGCCATGTGGACAGTGGCCAGGCTACAGATGCACTGGG 420
 DB 361 GCCGGGCTCTCACTCAAGTGGCCATGTGGACAGTGGCCAGGCTACAGATGCACTGGG 420

QY 421 AGCTGTGTCCAAAGCAGTGTCTAGCTGCGGGCCACTTCACTCTGCAGCAGCAGGGG 480
 DB 421 AGCTGTGTCCAAAGCAGTGTCTAGCTGCGGGCCACTTCACTCTGCAGCAGCAGGGG 480

QY 481 CAGGCCCCAGGGTGAGATGATCTGCCAGGCGTCTCGGGCAGGCCCACTATCAACCA 540
 DB 481 CAGGCCCCAGGGTGAGATGATCTGCCAGGCGTCTCGGGCAGGCCCACTATCAACCA 540

QY 541 GCCTGATCGGAAGATGGGAGGTCCACCTGCAGCAGACGACCATGCCAGGAGCGCTG 600
 DB 541 GCCTGATCGGAAGATGGGAGGTCCACCTGCAGCAGACGACCATGCCAGGAGCGCTG 600

QY 601 CCACTCTCTCTCTCTCCGAGCAGACATCGAGTGTCTGTGTCAGGCTCCAAACA 660
 DB 601 CCACTCTCTCTCTCTCCGAGCAGACATCGAGTGTCTGTGTCAGGCTCCAAACA 660

QY 661 ACGCCAATGTCCAGCAGCAGCCCTCAAGTGGTGGCCCGCCAGGTGGTCAAGAGATGG 720
 DB 661 ACGCCAATGTCCAGCAGCAGCCCTCAAGTGGTGGCCCGCCAGGTGGTCAAGAGATGG 720

QY 721 AGGACTGCGAGGTCCCTGAGAGCCCATCTTGGCTTCCCGTCTPACAGGAGCACC 780
 DB 721 AGGACTGCGAGGTCCCTGAGAGCCCATCTTGGCTTCCCGTCTPACAGGAGCACC 780

QY 781 GCCGCTCAGTGAAGAGAGTTTGGGGGTTTCAGATAGGAATGGGGAGTCTCAGGAC 840
 DB 781 GCCGCTCAGTGAAGAGAGTTTGGGGGTTTCAGATAGGAATGGGGAGTCTCAGGAC 840

QY 841 GCAAGCAGCAGCCATGTAGATGAACCTGCAGAGAGCCAGCAGCGCAGAGGACTGCA 900
 DB 841 GCAAGCAGCAGCCATGTAGATGAACCTGCAGAGAGCCAGCAGCGCAGAGGACTGCA 900

Db 841 GCAAGCAGCAGCCATGTAGATGAACCTGCAGAGAGCCAGCAGCGCAGAGGACTGCA 900
 QY 901 GGCCATCAGCGTGCACCTGTTCTGTTATTTGGAGTTTCATGCAAAATGAGTGTGTTTGTAGCTGC 960
 Db 901 GGCCATCAGCGTGCACCTGTTCTGTTATTTGGAGTTTCATGCAAAATGAGTGTGTTTGTAGCTGC 960
 QY 961 TCTTGGCCACAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAA 992
 Db 961 TCTTGGCCACAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAAATAAAAAA 992

RESULT 3
 ABK33610
 ID ABK33610 standard; cDNA; 992 BP.
 XX
 AC ABK33610;
 XX
 DT 08-MAY-2002 (first entry)
 XX
 DE cDNA encoding human PRO protein, Seq ID No 149.
 XX
 KW Human; secreted protein; PRO; tumour; lung cancer; colon cancer;
 KW breast cancer; prostate tumour; rectal tumour; liver tumour;
 KW pericyte cell proliferation; chondrocyte cell proliferation;
 KW tumour necrosis factor-alpha; gene; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO200208288-A2.
 XX
 PD 31-JAN-2002.
 XX
 PF 29-JUN-2001; 2001WO-US21066.
 XX
 PR 20-JUL-2000; 2000US-219556P.
 PR 25-JUL-2000; 2000US-220585P.
 PR 25-JUL-2000; 2000US-220605P.
 PR 25-JUL-2000; 2000US-220607P.
 PR 25-JUL-2000; 2000US-220624P.
 PR 25-JUL-2000; 2000US-220638P.
 PR 25-JUL-2000; 2000US-220664P.
 PR 26-JUL-2000; 2000US-220893P.
 PR 28-JUL-2000; 2000WO-US20710.
 PR 23-AUG-2000; 2000WO-US23522.
 PR 24-AUG-2000; 2000WO-US23328.
 PR 15-SEP-2000; 2000US-000000P.
 PR 10-NOV-2000; 2000WO-US30873.
 PR 28-NOV-2000; 2000US-253646P.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 20-DEC-2000; 2000US-074259.
 PR 20-DEC-2000; 2000WO-US34956.
 PR 28-FEB-2001; 2001WO-US06520.
 PR 10-MAY-2001; 2001US-0854280.
 PR 25-MAY-2001; 2001WO-US17092.
 XX
 PA (GETH) GENENTECH INC.
 XX
 PI Baker KP, Desnoyers L, Gerritsen ME, Goddard A, Godowski PU;
 PI Grimaldi JC, Gurney AL, Smith V, Stephan JF, Watanabe CK, Wood WT;
 XX
 XX WPI; 2002-172001/22.
 DR P-PSDB; AAU83666.
 DR
 XX
 PT One hundred and twenty two nucleic acids encoding PRO polypeptides,
 PT useful for treating a PRO related disorder and for diagnosing tumours
 PT such as lung cancer, colon cancer, breast tumour, prostate tumour, rectal
 PT tumour or liver tumour -
 XX
 PS Claim 2; Figure 149; 359pp; English.
 XX
 CC The invention relates to one hundred and twenty two nucleic acids
 CC encoding PRO polypeptides. The sequences of the 122 PRO polynucleotides

CC encode human secreted proteins. The PRO nucleic acids, polypeptides,
CC agonists and antagonists are useful for treating a PRO related disorder.
CC The PRO polypeptides are useful for diagnosing tumours, especially lung
CC cancer, colon cancer, breast tumour, prostate tumour, rectal tumour or
CC liver tumour. The PRO polypeptides are useful for stimulating the
CC proliferation of, or gene expression, in pericyte cells, for stimulating
CC the proliferation or differentiation of chondrocyte cells, for
CC stimulating the release of tumour necrosis factor-alpha from human blood,
CC for stimulating or inhibiting the proliferation of normal human dermal
CC fibroblast cells. The PRO polypeptide may also be used as molecular
CC weight markers and for tissue typing. The PRO nucleic acids have
CC applications in molecular biology, including use as hybridisation probes,
CC and in chromosome and gene mapping. ABK33536-ABK33657 represent human
CC PRO protein coding sequences of the invention.
XX
SQ Sequence 992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;

Query Match 100.0%; Score 992; DB 24; Length 992;
Best Local Similarity 100.0%; Pred. No. 2.7e-236;
Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCACGAGCCAGCACTAGGAGGTTCTCACTCCGAGCAGAGGCCCTACCCACCAG 60
Db 1 GGCACGAGCCAGCACTAGGAGGTTCTCACTCCGAGCAGAGGCCCTACCCACCAG 60
QY 61 GCATGGGGCTCCCTGGGCTGTCTGCTGGCGTGGCTGCCAGCAGCTTCTCAAGG 120
Db 61 GCATGGGGCTCCCTGGGCTGTCTGCTGGCGTGGCTGCCAGCAGCTTCTCAAGG 120
QY 121 CACGGGAGGAGAAATTAACCCCTGTGCTCCATGTGCTCAAAAGTCTCTGGAAGTTTTC 180
Db 121 CACGGGAGGAGAAATTAACCCCTGTGCTCCATGTGCTCAAAAGTCTCTGGAAGTTTTC 180
QY 181 CCAAGGCCCGTGGTGCTCTATACCTGTGTGACACCCAGCAGCACCAGCCATCACT 240
Db 181 CCAAGGCCCGTGGTGCTCTATACCTGTGTGACACCCAGCAGCACCAGCCATCACT 240
QY 241 ATTCCCTCTCTGGAACCAAGAAATCAATCAAGTGGCAGAGAGGTGGTGAACCCAGCAGC 300
Db 241 ATTCCCTCTCTGGAACCAAGAAATCAATCAAGTGGCAGAGAGGTGGTGAACCCAGCAGC 300
QY 301 CGGCTCTCTTCAACCTCAACGTCACTCAAGTCCAGTCCAGACCTGCTCACCCTATTCT 360
Db 301 CGGCTCTCTTCAACCTCAACGTCACTCAAGTCCAGTCCAGACCTGCTCACCCTATTCT 360
QY 361 GCCGGGCGTCCCTCACTCAGGTGCCATGTGACAGTGCAGGCTACAGATGCATCTGGG 420
Db 361 GCCGGGCGTCCCTCACTCAGGTGCCATGTGACAGTGCAGGCTACAGATGCATCTGGG 420
QY 421 AGCTGTGGTCCAAAGCCAGTGTCTGAGCTGGGGCCAACTTCACTCTGAGCAGAGAGGGG 480
Db 421 AGCTGTGGTCCAAAGCCAGTGTCTGAGCTGGGGCCAACTTCACTCTGAGCAGAGAGGGG 480
QY 481 CAGGCCCGAGGTGGAGATGATCTGCAGCGCTCTCGGGCAGCCCACTTATCAACCA 540
Db 481 CAGGCCCGAGGTGGAGATGATCTGCAGCGCTCTCGGGCAGCCCACTTATCAACCA 540
QY 541 GCCTGATCGGGAAGGATGGGAGTCCACCTGAGCAGAGAGACCATGCACAGCAGCCTG 600
Db 541 GCCTGATCGGGAAGGATGGGAGTCCACCTGAGCAGAGAGACCATGCACAGCAGCCTG 600
QY 601 CCAACTTCTCTTCTCCGAGCAGACATCGGACTGGTTCTGGTCCAGGCTGCAACA 660
Db 601 CCAACTTCTCTTCTCCGAGCAGACATCGGACTGGTTCTGGTCCAGGCTGCAACA 660
QY 661 ACGCCAATGTCAGCAGAGCCCTTACAGTGTGCCCCAGGTGGTGACCAAGATGG 720
Db 661 ACGCCAATGTCAGCAGAGCCCTTACAGTGTGCCCCAGGTGGTGACCAAGATGG 720
QY 721 AGGACTGGAGGGTCCCTGGAGAGCCCATCTTGGCTTGGCTCTACAGAGACCC 780
Db 721 AGGACTGGAGGGTCCCTGGAGAGCCCATCTTGGCTTGGCTCTACAGAGACCC 780

QY 781 GCCGTCTGAGTGAAGAGAGCTTTGGGGGTTTCAGGATAGGAATGGGAGTTCAGAGAC 840
Db 781 GCCGTCTGAGTGAAGAGAGCTTTGGGGGTTTCAGGATAGGAATGGGAGTTCAGAGAC 840
QY 841 GCAAAGCAGCAGCCATAGTAATGAACCGTCCAGAGAGCCAAAGCAGCAGGACTGCA 900
Db 841 GCAAAGCAGCAGCCATAGTAATGAACCGTCCAGAGAGCCAAAGCAGCAGGACTGCA 900
QY 901 GGCCATCAGCGTGCACCTGTTTGGATTTTCATGCAAAATGAGTGTGTTTAGCTGC 960
Db 901 GGCCATCAGCGTGCACCTGTTTGGATTTTCATGCAAAATGAGTGTGTTTAGCTGC 960
QY 961 TCTTGGCCACAAAAAAAAAAAAAAAAAAAAAAAAAA 992
Db 961 TCTTGGCCACAAAAAAAAAAAAAAAAAAAAAAAAAA 992

RESULT 4
ABX80267
ID ABX80267 standard; DNA; 992 BP.
AC ABX80267;
XX
DT 28-APR-2003 (first entry)
XX
DE Novel human secreted or transmembrane protein PRO791 DNA.
XX
KW Human; PRO; hypertrophy of neonatal heart; angiogenesis; wound healing;
KW cardiac insufficiency disorder; cancer; tumour; immune response;
KW adrenal cortical capillary endothelial growth; c-fos induction;
KW vascular endothelial growth factor inhibition; VEGF inhibition;
KW endothelial cell growth inhibitor; T-lymphocytes stimulation;
KW retinal neurons cell survival; rod photoreceptor cell survival;
KW retinal disorder; retinitis pigmentosa; kidney disorder;
KW mammalian kidney mesangial cell proliferation; Berger disease;
KW dermatitis; herpeticiformis; Crohn's disease; chondrocyte proliferation;
KW chondrocyte redifferentiation; sports injury; arthritis; gene; ds.
XX
OS Homo sapiens.
XX
PN US2002132252-A1.
XX
PD 19-SEP-2002.
XX
PF 14-NOV-2001; 2001US-0990442.
XX
PR 05-NOV-1997; 97WO-US20069.
PR 16-SEP-1998; 98WO-US19330.
PR 17-SEP-1998; 98WO-US19437.
PR 07-OCT-1998; 98WO-US21141.
PR 01-DEC-1998; 98WO-US25108.
PR 05-JAN-1999; 99WO-US00106.
PR 08-MAR-1999; 99WO-US05028.
PR 15-SEP-1999; 99WO-US21090.
PR 30-NOV-1999; 99WO-US21547.
PR 01-DEC-1999; 99WO-US28313.
PR 01-DEC-1999; 99WO-US28301.
PR 16-DEC-1999; 99WO-US30095.
PR 20-DEC-1999; 99WO-US30911.
PR 06-JAN-2000; 2000WO-US00219.
PR 06-JAN-2000; 2000WO-US00376.
PR 11-FEB-2000; 2000WO-US03565.
PR 18-FEB-2000; 2000WO-US04341.
PR 22-FEB-2000; 2000WO-US04414.
PR 24-FEB-2000; 2000WO-US04914.
PR 24-FEB-2000; 2000WO-US05004.
PR 02-MAR-2000; 2000WO-US05841.
PR 10-MAR-2000; 2000WO-US06319.
PR 15-MAR-2000; 2000WO-US06884.
PR 20-MAR-2000; 2000WO-US07377.
PR 30-MAR-2000; 2000WO-US08439.

PR 15-MAY-2000; 2000WO-US13358.
 PR 17-MAY-2000; 2000WO-US13705.
 PR 22-MAY-2000; 2000WO-US14042.
 PR 30-MAY-2000; 2000WO-US14941.
 PR 02-JUN-2000; 2000WO-US15264.
 PR 28-JUL-2000; 2000WO-US20710.
 PR 11-AUG-2000; 2000WO-US20231.
 PR 23-AUG-2000; 2000WO-US23522.
 PR 24-AUG-2000; 2000WO-US23328.
 PR 08-NOV-2000; 2000WO-US30952.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 28-FEB-2001; 2001WO-US06520.
 PR 01-JUN-2001; 2001WO-US17800.
 PR 20-JUN-2001; 2001WO-US19692.
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 PR 18-JUN-1998; 98US-089907P.
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 PR 28-AUG-2001; 2001US-0941992.
 (GETH) GENENTECH INC.
 PI Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL;
 PI Ferrara N, Fong S, Gerber H, Gerritsen ME, Goddard A, Godowski PJ;
 PI Grimaldi JC, Gurney AL, Kijavlin IJ, Napier MA, Pan J, Paoni NP;
 PI Roy MA, Stewart TA, Tumas D, Watanabe CK, Williams PM, Wood WJ;

Zhang Z;
 WI; 2003-247083/24.
 P-ESDB; ABUS9108.
 Novel isolated PRO polypeptides e.g., PRO826, PRO1068, PRO1184, PRO1346 and PRO1375, which stimulate proliferation of stimulated T-lymphocytes are therapeutically useful for enhancing immune response and in cancer treatments
 Claim 2; Fig 152; 648pp; English.
 The invention describes an isolated human PRO polypeptide. The PRO polypeptides are useful in detecting PRO polypeptides in a sample, in linking a bioactive molecule to a cell expressing a PRO polypeptide, and in modulating at least one biological activity of a cell expressing a PRO polypeptide. PRO1312 stimulates hypertrophy of neonatal heart and is thus useful for treating cardiac insufficiency disorders. PRO1154 and PRO1186 stimulate adrenal cortical capillary endothelial growth, and PRO536, PRO943, PRO828, PRO1068 or PRO535, PRO826, PRO819, PRO1126, PRO1360 and PRO1387 induce c-fos in endothelial cells, and are thus useful for treating conditions or disorders where angiogenesis would be beneficial, e.g. wound healing and antagonist of this polypeptide are useful for treating cancerous tumours. PRO812 inhibits vascular endothelial growth factor (VEGF) stimulated proliferation of endothelial cells and is thus useful for inhibiting endothelial cell growth in mammals which would be beneficial in inhibiting tumour growth. PRO826, PRO1068, PRO1184, PRO1346 and PRO1375 stimulate proliferation of stimulated T-lymphocytes and are therapeutically useful for enhancing immune response. PRO828, PRO1068 or PRO1132 enhance survival of retinal neurons cells (PRO1132 is also enhances survival/proliferation of rod photoreceptor cells) and therefore are useful for treating retinal disorders of injuries, e.g. retinitis pigmentosa, AMD. PRO819, PRO813 and PRO1066 induce proliferation of mammalian kidney mesangial cells, and therefore are useful for treating kidney disorders associated with decreased mesangial cell function such as Berger disease or Crohn's nephropathies associated with dermatitis, herpetiformis or Crohn's disease. PRO1310, PRO844, PRO1312, PRO1192 and PRO1387 induce the proliferation and/or redifferentiation of chondrocytes in culture and are thus useful for treating sports injuries, and arthritis. This sequence represents a novel human PRO protein polynucleotide.
 Sequence 992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;

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Qy 181	CCAAAGCCCGTGGTGTCTATACCTGTGTGCACCCGAGCAGCACCACCCATCACT	240		
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Query Match 100.0%; Score 992; DB 25; Length 992;
Best Local Similarity 100.0%; Pred. No. 2,7e-236;
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Qy 121 CACGGGAGGAGAAATTTACCCCTGTGTCTCCATTTGCTCAAAAGTCTCGAAAGTTTTC 180
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XX AC ABX81154;
XX DT 22-APR-2003 (first entry)
XX DE Novel human secreted or transmembrane protein PRO791 DNA.
XX KW Human; PRO; hypertrophy of neonatal heart; angiogenesis; wound healing;
KW cardiac insufficiency disorder; cancer; tumour; immune response;
KW adrenal cortical capillary endothelial growth; c-fos induction;
KW vascular endothelial growth factor inhibition; VEGF inhibition;
KW endothelial cell growth inhibitor; T-lymphocytes stimulation;
KW retinal neurons cell survival; rod photoreceptor cell survival;
KW retinal disorder; retinitis pigmentosa; kidney disorder;
KW mammalian kidney mesangial cell proliferation; Berger disease;
KW dermatitis; herpeticiformis; Crohn's disease; chondrocyte proliferation;
KW chondrocyte redifferentiation; sports injury; arthritis; gene; ds.
XX OS Homo sapiens.
XX PN US2003027985-A1.
XX PD 06-FEB-2003.
XX PF 14-NOV-2001; 2001US-0990562.
XX QY 05-NOV-1997; 97WO-US200069.
PR 16-SEP-1998; 98WO-US19330.
PR 17-SEP-1998; 98WO-US19437.
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PR 26-AUG-1998; 98US-098014P.

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Db 721 AGGACTGGCAGGGTCCCTGGAGAGCCCATCTTGTGCTTGGCTTACAGAGCAACC 780
Qy 781 GCCGCTCTGAGTGAAGAGGAGTTTGGGGGGTTTCCAGGATAGGAATGGGAGGTGAGGAC 840
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Qy 841 GCAAAGCAGCAGCCATGTAGATGAACCGTCCAGAGAGCCACGACCGGACAGGAGTGC 900
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CC encoding them. The polypeptides can be used to raise antibodies that
CC specifically bind to the PRO polypeptide, for linking a bioactive
CC molecule to a cell expressing a PRO protein and for modulating at least
CC one biological activity of a cell. The PRO polypeptides or
CC polynucleotides are also useful in gene therapy, in chromosome
CC identification, as chromosome markers, or in generating probes. The PRO
CC polypeptides are useful as molecular markers for protein
CC electrophoresis, and the isolated nucleic acids may be used for
CC recombinantly expressing those markers. The PRO polypeptides and nucleic
CC acids may also be used in tissue typing. Anti-PRO antibodies are useful
CC in diagnostic assays for PRO, and in affinity purification of PRO from
CC recombinant cell culture or natural sources. The sequences presented in
CC ABX90083-ABX90468 are the genes encoding, the primers amplifying and the
CC probes detecting the PRO polynucleotides of the invention.
CC Note: The sequence data for this patent is also available in electronic
CC format from USPTO at seqdata.uspto.gov/sequence.html.
XX

SQ Sequence 992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;

Query Match 100.0%; Score 992; DB 25; Length 992;
Best Local Similarity 100.0%; Pred. No. 2.7e-236;
Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;
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XX AC Human PRO polynucleotide #61.
XX DT 14-APR-2003 (first entry)
XX DE Human PRO polynucleotide #61.
XX KW Human; PRO; gene; ss; cytostatic; tumour; cancer; breast; lung; stomach;
KW liver; horse; cow; dog; cat; sheep; pig; goat; rabbit; ADEPT;
KW antibody-dependent enzyme mediated prodrug therapy.
XX OS Homo sapiens.
XX PN US2003027163-A1.
XX PD 06-FEB-2003.
XX PF 15-NOV-2001; 2001US-0997666.
XX PR 05-NOV-1997; 97NO-US20069.
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Best Local Similarity 100.0%; Pred. No. 2.7e-236;
Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

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XX colon cancer; lung cancer; breast cancer; cancer; gene therapy.
XX Homo sapiens.
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XX 03-OCT-2002.
XX 19-NOV-2001; 2001US-0989721.
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PR 10-JUN-1998; 98US-088824P.
PR 10-JUN-1998; 98US-088826P.
PR 11-JUN-1998; 98US-088858P.
PR 11-JUN-1998; 98US-088861P.
PR 11-JUN-1998; 98US-088876P.
PR 12-JUN-1998; 98US-089105P.
PR 16-JUN-1998; 98US-089440P.
PR 16-JUN-1998; 98US-089512P.
PR 16-JUN-1998; 98US-089514P.
PR 17-JUN-1998; 98US-089532P.
PR 17-JUN-1998; 98US-089538P.
PR 17-JUN-1998; 98US-089598P.
PR 17-JUN-1998; 98US-089599P.
PR 17-JUN-1998; 98US-089600P.
PR 17-JUN-1998; 98US-089653P.
PR 18-JUN-1998; 98US-089801P.
PR 18-JUN-1998; 98US-089907P.
PR 18-JUN-1998; 98US-089908P.
PR 28-AUG-2001; 2001US-0941992.
XX (GETH) GENENTECH INC.
XX
XX Ashkenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL,
PI Ferrara N, Fong S, Gerber H, Gerritsen ME, Goddard A, Godowski PJ,
PI Grimaldi JC, Gurney AL, Kijavini IJ, Napier MA, Pan J, Paoni NF,
PI Roy MA, Stewart TA, Tumas D, Watanabe CK, Williams FM, Wood WI,
PI Zhang Z;
XX
XX WPI; 2003-155950/15.
DR P-PSDB; ABUS8960.
XX
XX
PT New secreted and transmembrane PRO polypeptides (e.g. PRO183, PRO184,
PT PRO361 or PRO846) useful as targets for therapeutic intervention in
PT cancers (e.g. lung or breast cancers), or for diagnosing these cancers
PT
XX
XX Claim 2; Fig 150; 647pp; English.
XX
XX The invention discloses isolated PRO secreted/transmembrane polypeptides
CC comprising a sequence without signal peptide and the nucleic acid
CC encoding them. The polypeptides can be used to raise antibodies that
CC specifically bind to the PRO polypeptide, for linking a bioactive
CC molecule to a cell expressing a PRO protein and for modulating at least
CC one biological activity of a cell. The PRO polypeptides or
CC polynucleotides are also useful as pharmaceuticals, diagnostics,
CC biosensors or bioeffectors, for detecting or treating e.g. tumours in
CC mammals, e.g. humans, dogs, cats, cattle, horses, sheep, pigs, goats or
CC rabbits as targets for therapeutic intervention in certain cancers (e.g.

CC colon, lung or breast cancers) and diagnostic determination of the
CC presence of these cancers. The PRO polypeptides are also useful as
CC molecular weight markers or for chromosome identification. The PRO genes
CC are useful as hybridisation probes or for screening libraries of human
CC cDNA, genomic DNA or mRNA. The PRO genes may also be used in gene
CC therapy, particularly for replacing a defective gene. The sequences
CC presented in ABX79290-ABX79675 are the genes encoding, the primers
CC amplifying and the probes detecting the PRO polynucleotides of the
CC invention.
CC Note: The sequence data for this patent is also available in electronic
CC format from USPTO at seqdata.uspto.gov/sequence.html.
XX

SQ Sequence 992 BP; 235 A; 301 C; 282 G; 174 T; 0 other;

Query Match 100.0%; Score 992; DB 25; Length 992;
Best Local Similarity 100.0%; Pred. No. 2,7e-236;
Matches 992; Conservative 0; Mismatches 0; Indels 0; Gaps 0;

QY 1 GGCACGAGCCAGGAAC TAGAGGTTCTCACTGCCGAGCAGAGGCCCTACACCCACCGAG 60
Db 1 GGCACGAGCCAGGAAC TAGAGGTTCTCACTGCCGAGCAGAGGCCCTACACCCACCGAG 60
QY 61 GCATGGGGCTCCCTGGGCTGTTCTGCTGGCCGCTGCTGGCTGCACAGGTTTCCAAG 120
Db 61 GCATGGGGCTCCCTGGGCTGTTCTGCTGGCCGCTGCTGGCTGCACAGGTTTCCAAG 120
QY 121 CACGGGAGGAAGAAATTACCCCTGTGGTCTCCATTGCCCTACAAAGTCCTGGAAGTTTTC 180
Db 121 CACGGGAGGAAGAAATTACCCCTGTGGTCTCCATTGCCCTACAAAGTCCTGGAAGTTTTC 180
QY 181 CCAAGGCCGCTGGGTGCTCATAAAGTGTGTCACCCCGAGCCACCCGCGCCATCACT 240
Db 181 CCAAGGCCGCTGGGTGCTCATAAAGTGTGTCACCCCGAGCCACCCGCGCCATCACT 240
QY 241 ATTCCCTCTGTGGAACCAAGACATCAAGTGGCCCAAGAGTGGTGAAGACCCAGAGC 300
Db 241 ATTCCCTCTGTGGAACCAAGACATCAAGTGGCCCAAGAGTGGTGAAGACCCAGAGC 300
QY 301 CGGCTCTCTTCAACCTCAACGTCACACTCAAGTCCAGCTCCAGCTGCTCACTACTTCT 360
Db 301 CGGCTCTCTTCAACCTCAACGTCACACTCAAGTCCAGCTCCAGCTGCTCACTACTTCT 360
QY 361 GCGGGGCTCTCCACCTCAGTGCCCATGTGGAACAGTGCACAGGTACAGATGCACTGGG 420
Db 361 GCGGGGCTCTCCACCTCAGTGCCCATGTGGAACAGTGCACAGGTACAGATGCACTGGG 420
QY 421 AGCTGTGTCCAAAGCCAGTGTCTGAGCTGGGGCCCACTTCACTCTGAGGACAGAGGG 480
Db 421 AGCTGTGTCCAAAGCCAGTGTCTGAGCTGGGGCCCACTTCACTCTGAGGACAGAGGG 480
QY 481 CAGGCCCCAGGGTGGAGATGATCTGCCAGGGGCTCTCGGGGAGGCCCACTATCAACA 540
Db 481 CAGGCCCCAGGGTGGAGATGATCTGCCAGGGGCTCTCGGGGAGGCCCACTATCAACA 540
QY 541 GCCTGATCGGAGGATGGGAGGTCCACCTGACAGCAGACAGACCATGCCAGGAGGCTG 600
Db 541 GCCTGATCGGAGGATGGGAGGTCCACCTGACAGCAGACAGACCATGCCAGGAGGCTG 600
QY 601 CCAACTTCTCCTTCTGCGGAGCCAGACATCGGACTGTTCTGTGGTGCAGGCTGCAACA 660
Db 601 CCAACTTCTCCTTCTGCGGAGCCAGACATCGGACTGTTCTGTGGTGCAGGCTGCAACA 660
QY 661 AGCCCAAGTCCAGCAGACAGCCCTTCAAGTGTGTGCCCGAGGTGGTACAGAGATGG 720
Db 661 AGCCCAAGTCCAGCAGACAGCCCTTCAAGTGTGTGCCCGAGGTGGTACAGAGATGG 720
QY 721 AGGACTGCGAGGTTCCTCTGAGAGCCCATCTTTCCTTGCCTTCCGCTCTACAGGAGCAC 780
Db 721 AGGACTGCGAGGTTCCTCTGAGAGCCCATCTTTCCTTGCCTTCCGCTCTACAGGAGCAC 780
QY 781 GCGTGTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGGATAGGGAATGGGAGGTTCAGAGGAC 840
Db 781 GCGTGTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGGATAGGGAATGGGAGGTTCAGAGGAC 840

QY 841 GCAAAGCAGCAGCCATGTAGTAATGAACCTCCAGAGAGCCAGCAGCGGACGAGGACTGCA 900
 |||
 Db 841 GCAAGAGCAGCAGCCATGTAGTAATGAACCTCCAGAGAGCCAGCAGCGGACGAGGACTGCA 900
 |||
 QY 901 GGCATCAGCGTGCATGTTCGTATTTGGAGTTCATGCAAAATGATGTGTTTTAGCTGC 960
 |||
 Db 901 GGCATCAGCGTGCATGTTCGTATTTGGAGTTCATGCAAAATGATGTGTTTTAGCTGC 960
 |||
 QY 961 TCTTGCACAAAAAAAAAAAAAAAAAAAAAAAAA 992
 |||
 Db 961 TCTTGCACAAAAAAAAAAAAAAAAAAAAAAAAA 992
 |||

RESULT 10

ABX64090

ID ABX64090 standard; cDNA; 992 BP.

XX AC ABX64090;

XX 26-FEB-2003 (first entry)

XX cDNA encoding human PRO809 polypeptide.

XX Human; PRO polypeptide; secreted protein; transmembrane protein;
 KW genetic disorder; antibacterial; immunosuppressive; transgenic;
 KW gene therapy; gene; ss.

XX OS Homo sapiens.

XX PN US2002103125-A1.

XX PD 01-AUG-2002.

XX 20-NOV-2001; 2001US-0989731.

XX 05-NOV-1997; 97WO-US20069.

PR 16-SEP-1998; 98WO-US19330.

PR 17-SEP-1998; 98WO-US19437.

PR 07-OCT-1998; 98WO-US21141.

PR 01-DEC-1998; 98WO-US25108.

PR 05-JAN-1999; 99WO-US00106.

PR 08-MAR-1999; 99WO-US05028.

PR 02-JUN-1999; 99WO-US12252.

PR 15-SEP-1999; 99WO-US21090.

PR 15-SEP-1999; 99WO-US21547.

PR 30-NOV-1999; 99WO-US28313.

PR 01-DEC-1999; 99WO-US28301.

PR 01-DEC-1999; 99WO-US28634.

PR 16-DEC-1999; 99WO-US30095.

PR 20-DEC-1999; 99WO-US30911.

PR 06-JAN-2000; 2000WO-US00219.

PR 06-JAN-2000; 2000WO-US00376.

PR 11-FEB-2000; 2000WO-US03565.

PR 18-FEB-2000; 2000WO-US04341.

PR 22-FEB-2000; 2000WO-US04414.

PR 24-FEB-2000; 2000WO-US04914.

PR 24-FEB-2000; 2000WO-US05004.

PR 10-MAR-2000; 2000WO-US05841.
 PR 15-MAR-2000; 2000WO-US06319.
 PR 20-MAR-2000; 2000WO-US06884.
 PR 30-MAR-2000; 2000WO-US07377.
 PR 15-MAY-2000; 2000WO-US08439.
 PR 17-MAY-2000; 2000WO-US13358.
 PR 22-MAY-2000; 2000WO-US13705.
 PR 30-MAY-2000; 2000WO-US14042.
 PR 02-JUN-2000; 2000WO-US14941.
 PR 28-JUL-2000; 2000WO-US20710.
 PR 11-AUG-2000; 2000WO-US22031.
 PR 23-AUG-2000; 2000WO-US23522.
 PR 24-AUG-2000; 2000WO-US23328.
 PR 08-NOV-2000; 2000WO-US30952.
 PR 01-DEC-2000; 2000WO-US32678.
 PR 28-FEB-2001; 2001WO-US06520.
 PR 01-JUN-2001; 2001WO-US17800.
 PR 20-JUN-2001; 2001WO-US19692.
 PR 29-JUN-2001; 2001WO-US21066.
 PR 09-JUL-2001; 2001WO-US21735.
 PR 16-JUN-1997; 97US-049787P.
 PR 17-OCT-1997; 97US-062250P.
 PR 12-NOV-1997; 97US-065186P.
 PR 13-NOV-1997; 97US-065311P.
 PR 24-NOV-1997; 97US-066770P.
 PR 25-FEB-1998; 98US-075945P.
 PR 20-MAR-1998; 98US-078910P.
 PR 28-APR-1998; 98US-083322P.
 PR 07-MAY-1998; 98US-084600P.
 PR 28-MAY-1998; 98US-087106P.
 PR 02-JUN-1998; 98US-087607P.
 PR 02-JUN-1998; 98US-087609P.
 PR 02-JUN-1998; 98US-087759P.
 PR 03-JUN-1998; 98US-087827P.
 PR 04-JUN-1998; 98US-088021P.
 PR 04-JUN-1998; 98US-088025P.
 PR 04-JUN-1998; 98US-088026P.
 PR 04-JUN-1998; 98US-088028P.
 PR 04-JUN-1998; 98US-088029P.
 PR 04-JUN-1998; 98US-088030P.
 PR 04-JUN-1998; 98US-088033P.
 PR 04-JUN-1998; 98US-088326P.
 PR 05-JUN-1998; 98US-088167P.
 PR 05-JUN-1998; 98US-088202P.
 PR 05-JUN-1998; 98US-088212P.
 PR 05-JUN-1998; 98US-088217P.
 PR 09-JUN-1998; 98US-088655P.
 PR 10-JUN-1998; 98US-088734P.
 PR 10-JUN-1998; 98US-088738P.
 PR 10-JUN-1998; 98US-088742P.
 PR 10-JUN-1998; 98US-088810P.
 PR 10-JUN-1998; 98US-088824P.
 PR 11-JUN-1998; 98US-088858P.
 PR 11-JUN-1998; 98US-088861P.
 PR 11-JUN-1998; 98US-088876P.
 PR 12-JUN-1998; 98US-089105P.
 PR 16-JUN-1998; 98US-089440P.
 PR 16-JUN-1998; 98US-089512P.
 PR 17-JUN-1998; 98US-089514P.
 PR 17-JUN-1998; 98US-089532P.
 PR 17-JUN-1998; 98US-089538P.
 PR 17-JUN-1998; 98US-089598P.
 PR 17-JUN-1998; 98US-089599P.
 PR 17-JUN-1998; 98US-089600P.
 PR 18-JUN-1998; 98US-089653P.
 PR 18-JUN-1998; 98US-089801P.
 PR 18-JUN-1998; 98US-089907P.
 PR 18-JUN-1998; 98US-089908P.
 PR 28-AUG-2001; 2001US-0941992.

(GETH) GENENTECH LTD.

Askenazi AJ, Baker KP, Botstein D, Desnoyers L, Eaton DL;
 Ferrara N, Fong S, Gerber H, Gerritsen ME, Goddard A, Godowski P;
 Grimaldi JC, Gurney AL, Kljavin IO, Napier MA, Pan J, Paoni NF;
 Roy MA, Stewart TA, Tumas D, Watanabe CK, Williams PM, Wood WI;
 Zhang Z;

WPI: 2003-102117/09.

P-PSDB; ABU13920.

Novel secreted and transmembrane polypeptide for modulating biological
 activity of cell expressing the polypeptide, identifying agonists or
 antagonists of polypeptide, and as molecular weight markers -

Claim 2; Fig 150; 649pp; English.


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Db 481 CAGCCCCAGGTTGGAGATGATCTGCCAGCGCTCCTCGGCGAGCCCCACCTATCACCACA 540
QY 541 GCCTGATCGGAAGGATGGCAGGTTCCACTCTCAGCAGAGACCATGTCACAGGCGAGCCTG 600
Db 541 GCCTGATCGGAAGGATGGCAGGTTCCACTCTCAGCAGAGACCATGTCACAGGCGAGCCTG 600
QY 601 CCAACTTCTCTTCTCTCTCCAGCAGCAGCATCGGACTGGTTCTGGTGCAGGCTGCAACA 660
Db 601 CCAACTTCTCTTCTCTCTCCAGCAGCAGCATCGGACTGGTTCTGGTGCAGGCTGCAACA 660
QY 661 ACGCCAATGTCAGCAGCAGCGCCTCACAGTGTGCCCGCAGTGTGACCAAGATGG 720
Db 661 ACGCCAATGTCAGCAGCAGCGCCTCACAGTGTGCCCGCAGTGTGACCAAGATGG 720
QY 721 AGCACTGGCAGGTTCCCTCGAGAGGCCCACTCTTGCCTTGCCTTCCGCTCTACAGGAGCACCC 780
Db 721 AGCACTGGCAGGTTCCCTCGAGAGGCCCACTCTTGCCTTGCCTTCCGCTCTACAGGAGCACCC 780
QY 781 GCCGTCTGATGAAGAGGATTTGGGGGTTTCAGGATAGGAAATGGGAGGTTCAGAGGAC 840
Db 781 GCCGTCTGATGAAGAGGATTTGGGGGTTTCAGGATAGGAAATGGGAGGTTCAGAGGAC 840
QY 841 GCAAGCAGCAGCATGTAGAATGAACCGTCCAGAGAGCCCAAGCAGGAGGACTGCA 900
Db 841 GCAAGCAGCAGCATGTAGAATGAACCGTCCAGAGAGCCCAAGCAGGAGGACTGCA 900
QY 901 GGCCATCAGCGTCACTGTTCTGATTTGGAGTTTCATGCAAAATGAGTGTGTTTAGCTGC 960
Db 901 GGCCATCAGCGTCACTGTTCTGATTTGGAGTTTCATGCAAAATGAGTGTGTTTAGCTGC 960
QY 961 TCTTGCACAAAAAATAAAAAAAAAAAAAAAAAAAAAA 992
Db 961 TCTTGCACAAAAAATAAAAAAAAAAAAAAAAAAAAAA 992

RESULT 12
AAZ65030
ID AAZ65030 standard; cDNA; 991 BP.
XX
AC AAZ65030;
XX
DT 05-APR-2000 (first entry)
XX
DE Membrane-bound protein PRO809 encoding cDNA.
XX
KW Membrane-bound polypeptide; PRO polypeptide; LDL receptor; TIE ligand;
KW pharmaceutical; receptor immunoadhesin; gene mapping; ss.
XX
OS Homo sapiens.
XX
PN WO963088-A2.
XX
PD 09-DEC-1999.
XX
PF 02-JUN-1999; 99WO-US12252.
XX
PR 02-JUN-1998; 98US-0087607.
PR 02-JUN-1998; 98US-0087609.
PR 02-JUN-1998; 98US-0087759.
PR 03-JUN-1998; 98US-0087827.
PR 04-JUN-1998; 98US-0088021.
PR 04-JUN-1998; 98US-0088025.
PR 04-JUN-1998; 98US-0088028.
PR 04-JUN-1998; 98US-0088029.
PR 04-JUN-1998; 98US-0088030.
PR 04-JUN-1998; 98US-0088033.
PR 04-JUN-1998; 98US-0088326.
PR 05-JUN-1998; 98US-0088167.
PR 05-JUN-1998; 98US-0088202.
PR 05-JUN-1998; 98US-0088212.
PR 05-JUN-1998; 98US-0088217.
PR 09-JUN-1998; 98US-0088655.
PR 10-JUN-1998; 98US-0088722.
PR 10-JUN-1998; 98US-0088730.
PR 10-JUN-1998; 98US-0088734.
PR 10-JUN-1998; 98US-0088738.
PR 10-JUN-1998; 98US-0088740.
PR 10-JUN-1998; 98US-0088741.
PR 10-JUN-1998; 98US-0088742.
PR 10-JUN-1998; 98US-0088810.
PR 10-JUN-1998; 98US-0088811.
PR 10-JUN-1998; 98US-0088824.
PR 10-JUN-1998; 98US-0088825.
PR 10-JUN-1998; 98US-0088826.
PR 11-JUN-1998; 98US-0088858.
PR 11-JUN-1998; 98US-0088861.
PR 11-JUN-1998; 98US-0088863.
PR 11-JUN-1998; 98US-0088876.
PR 12-JUN-1998; 98US-0089090.
PR 12-JUN-1998; 98US-0089105.
PR 16-JUN-1998; 98US-0089440.
PR 16-JUN-1998; 98US-0089512.
PR 16-JUN-1998; 98US-0089514.
PR 17-JUN-1998; 98US-0089532.
PR 17-JUN-1998; 98US-0089538.
PR 17-JUN-1998; 98US-0089598.
PR 17-JUN-1998; 98US-0089599.
PR 17-JUN-1998; 98US-0089600.
PR 17-JUN-1998; 98US-0089653.
PR 18-JUN-1998; 98US-0089801.
PR 18-JUN-1998; 98US-0089907.
PR 18-JUN-1998; 98US-0089908.
PR 19-JUN-1998; 98US-0089947.
PR 19-JUN-1998; 98US-0089948.
PR 19-JUN-1998; 98US-0089952.
PR 22-JUN-1998; 98US-0090246.
PR 22-JUN-1998; 98US-0090252.
PR 22-JUN-1998; 98US-0090254.
PR 23-JUN-1998; 98US-0090349.
PR 23-JUN-1998; 98US-0090355.
PR 24-JUN-1998; 98US-0090429.
PR 24-JUN-1998; 98US-0090431.
PR 24-JUN-1998; 98US-0090435.
PR 24-JUN-1998; 98US-0090444.
PR 24-JUN-1998; 98US-0090445.
PR 24-JUN-1998; 98US-0090461.
PR 24-JUN-1998; 98US-0090472.
PR 24-JUN-1998; 98US-0090535.
PR 24-JUN-1998; 98US-0090538.
PR 24-JUN-1998; 98US-0090540.
PR 24-JUN-1998; 98US-0090557.
PR 25-JUN-1998; 98US-0090676.
PR 25-JUN-1998; 98US-0090678.
PR 25-JUN-1998; 98US-0090688.
PR 25-JUN-1998; 98US-0090690.
PR 25-JUN-1998; 98US-0090691.
PR 25-JUN-1998; 98US-0090694.
PR 25-JUN-1998; 98US-0090695.
PR 26-JUN-1998; 98US-0090696.
PR 26-JUN-1998; 98US-0090862.
PR 26-JUN-1998; 98US-0090863.
PR 01-JUL-1998; 98US-0091358.
PR 01-JUL-1998; 98US-0091360.
PR 02-JUL-1998; 98US-0091478.
PR 02-JUL-1998; 98US-0091486.
PR 02-JUL-1998; 98US-0091519.
PR 02-JUL-1998; 98US-0091626.
PR 02-JUL-1998; 98US-0091628.
PR 02-JUL-1998; 98US-0091633.
PR 02-JUL-1998; 98US-0091646.
PR 02-JUL-1998; 98US-0091673.
PR 07-JUL-1998; 98US-0091978.
PR 07-JUL-1998; 98US-0091982.
PR 09-JUL-1998; 98US-0092182.
PR 10-JUL-1998; 98US-0092472.
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Db 961 TCTTGCCACAAAAA 991

RESULT 13
AAH64779
AAH64779 standard; cDNA; 1047 BP.
AC
AAH64779;
11-SEP-2001 (first entry)
DE Human secreted protein cDNA, SEQ ID NO: 55.
XX
KW Human; secreted protein; gene therapy; vaccine; treatment; diagnosis;
KW GENSET; ss.
XX
OS Homo sapiens.
XX
PN WO200142451-A2.
XX
PD 14-JUN-2001.
XX
XX 07-DEC-2000; 2000WO-IB01938.
XX
XX 08-DEC-1999; 99US-0169629.
PR 06-MAR-2000; 2000US-0187470.
XX
XX (GIST) GENSET.
XX
PI Dumas Milne Edwards J, Bougueleret L, Jobert S;
XX
XX WPI; 2001-367870/38.
DR P-PSDB; AAG89176.
XX
PT Full length GENSET human nucleic acids encoding potentially secreted
PT proteins, useful in gene therapy and vaccination against a variety of
PT diseases, and for diagnosis of those diseases -
XX
XX Claim 7; Page 620-621; 921pp; English.
XX
CC The invention relates to full length GENSET human nucleic acids encoding
CC potentially secreted proteins. The nucleic acids and the polypeptides
CC they encode may be used in the prevention, treatment and diagnosis of
CC diseases associated with inappropriate GENSET gene expression. For
CC example, they be used to treat disorders associated with decreased
CC GENSET gene expression by rectifying mutations or deletions in a
CC patient's genome that affect the activity of GENSET or by supplementing
CC the patient's own production of GENSET polypeptides. Conversely,
CC antisense nucleic acid molecules may be administered to down regulate
CC GENSET expression by binding with the cells' own genes and preventing
CC their expression. The sense and antisense nucleic acids may also be
CC used as DNA probes in diagnostic assays to detect and quantitate the
CC presence of similar nucleic acid sequences in samples, and hence to
CC determine which patients may be in need of restorative therapy.
CC The GENSET polypeptides may be used as antigens in the production of
CC antibodies and in assays to identify modulators (agonists and
CC antagonists) of GENSET polypeptide expression and activity. The
CC present sequence is a GENSET nucleic acid of the invention.
XX
SQ Sequence 1047 BP; 235 A; 324 C; 302 G; 186 T; 0 other;

Query Match 85.0%; Score 843; DB 22; Length 1047;
Best Local Similarity 90.7%; Pred. No. 2.6e-199;
Matches 950; Conservative 0; Mismatches 0; Indels 97; Gaps 1;

QY 37 AGCAGAGCCCTACACCCACCGAGGATGGGCTCCCTGGGCTGTCTGTTGGCCGTGC 96
DB 1 AGCAGAGCCCTACACCCACCGAGGATGGGCTCCCTGGGCTGTCTGTTGGCCGTGC 60
QY 97 TGGCTGCCAGCAGCTTCTCCAGAGCAGCGGAGAGAAATACCCCTGTGTTCCATTG 156
DB 61 TGGCTGCCAGCAGCTTCTCCAGAGCAGCGGAGAGAAATACCCCTGTGTTCCATTG 120

QY 157 CCTACAAAGTCTCTGGAAGTTTCCCAAGGCGCTGGGTGCTCATAACTGTGTGCAC 216
DB 121 CCTACAAAGTCTCTGGAAGTTTCCCAAGGCGCTGGGTGCTCATAACTGTGTGCAC 180
QY 217 CCCAGCCACACCGCCCATCACCATTATTCCTCTGTGTGAACCAAGAACATCAAGTGGCCA 276
DB 181 CCCAGCCACACCGCCCATCACCATTATTCCTCTGTGTGAACCAAGAACATCAAGTGGCCA 240
QY 277 AGAAGTGTGTGAAGACCCACGAGCCGCTCTTCAACTCAACCTCAACCTCAACCTCAAGTCCA 336
DB 241 AGAAGTGTGTGAAGACCCACGAGCCGCTCTTCAACTCAACCTCAACCTCAACCTCAAGTCCA 300
QY 337 GTCCACAGCTGTCACTACTTCTGCGGGCGCTCTCCACCTCAGTGGCCATGTGGACA 396
DB 301 GTCCACAGCTGTCACTACTTCTGCGGGCGCTCTCCACCTCAGTGGCCATGTGGACA 360
QY 397 GTGCCAGGCTTACAGATGCATCTGGGAGCTGTGTCTCAAGCCAGTGTCTGAGCTGCGGGCCA 456
DB 361 GTGCCAGGCTTACAGATGCATCTGGGAGCTGTGTCTCAAGCCAGTGTCTGAGCTGCGGGCCA 420
QY 457 ACTTCACTCTGAGGACAGAGGGGCGAGGCGCCAGGCTGAGATGATCTGCCAGGCGCTCT 516
DB 421 ACTTCACTCTGAGGACAGAGGGGCGAGGCGCCAGGCTGAGATGATCTGCCAGGCGCTCT 480
QY 517 CGGGCAGCCCACTATCACCACAGCTGATCGGGAAGATGGGAGCTCCACCTGCAGC 576
DB 481 CGGGCAGCCCACTATCACCACAGCTGATCGGGAAGATGGGAGCTCCACCTGCAGC 540
QY 577 AGAGACCATGCCACAGGAGCTTGCACACTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 636
DB 541 AGAGACCATGCCACAGGAGCTTGCACACTTCTCTCTCTCTCTCTCTCTCTCTCTCTCTCT 600
QY 637 GGTTCGTGTGCGAGCTGCAACACAGCCCAATGTTCAGCACAGGCGCCCTCACAGTGGTGC 696
DB 601 GGTTCGTGTGCGAGCTGCAACACAGCCCAATGTTCAGCACAGGCGCCCTCACAGTGGTGC 660
QY 697 CC----- 698
DB 661 CCCAGAGGGTTGCCAGGGGCAACCCACCATCGTGTGGTGGAGCTTGGCTCCACTG 720
QY 699 -----CCAGGTGGTGACCAAGATG 719
DB 721 CGGCCATCACCTCCAGGATGTGGGCTGGACCACTGGGSCCAGGTGTGACCAAGATG 780
QY 720 GAGGACTGCGAGGTCCCTTGAGAGGCCCATCTCTTGGCTTCCCGCTCTACAGGAGCAC 779
DB 781 GAGGACTGCGAGGTCCCTTGAGAGGCCCATCTCTTGGCTTCCCGCTCTACAGGAGCAC 840
QY 780 CGCCCTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGATAGGGAATGGGAGGTTCAGAGA 839
DB 841 CGCCCTCTGAGTGAAGAGGAGTTTGGGGGTTTCAGATAGGGAATGGGAGGTTCAGAGA 900
QY 840 CGCAAGCAGCAGCCATCTAGATGAACCGTCCAGAGAGCCCAAGCACCGCAGAGCACTGC 899
DB 901 CGCAAGCAGCAGCCATCTAGATGAACCGTCCAGAGAGCCCAAGCACCGCAGAGCACTGC 960
QY 900 AGGCCATCAGGTGCACTGTCTGATTTTGGAGTTTCATGCAAAATGAGTGTGTTAGCTG 959
DB 961 AGGCCATCAGGTGCACTGTCTGATTTTGGAGTTTCATGCAAAATGAGTGTGTTAGCTG 1020
QY 960 CTCCTGGCCACAAAAA 986
DB 1021 CTCCTGGCCACAAAAA 1047

RESULT 14
AAH99131
AAH99131 standard; cDNA; 935 BP.
XX
AC AAH99131;
XX
DT 12-OCT-2001 (first entry)
XX

(HUMA-) HUMAN GENOME SCI INC.

Rosen GA, Ruben SM, Komatsoulis G;

WFI; 2000-579483/54.

P-PSDB; AAB39216.

Isolated nucleic acid molecule encoding a human secreted protein is used in preventing, treating or ameliorating a medical condition -

Claim 1; Page 363-364; 434pp; English.

The polynucleotide sequences given in AAC74223-C74279 encode the human secreted proteins represented in AAB39179-B39226. Sequences AAB39227-B39308 are alternative proteins encoded by the genes, and also protein sequences with which they share homology. The proteins have activities based on the tissues and cells in which they are expressed. Examples of activities include: immunosuppressive; antiarthritic; antirheumatic; antiproliferative; cytostatic; cardiac; vasotropic; cerebroprotective; neurotropic; neuroprotective; antibacterial; virucide; fungicide; and opthalmological. The human secreted proteins, polynucleotides, antagonists and agonists of the invention may be useful in the treatment, prevention, and/or diagnosis of various disease, disorders and conditions such as autoimmune diseases e.g. rheumatoid arthritis, hyperproliferative disorders e.g. neoplasms of the breast or liver, cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g. Alzheimer's disease, infections caused by bacteria, viruses and fungi and ocular disorders e.g. corneal infection. The polypeptides can also be used to aid wound healing and epithelial cell proliferation, to regenerate tissues, maintain organs before transplantation, in chemotaxis and as a food additive or preservative e.g. to increase storage capabilities. Sequences AAC74214-C74222 and AAB39178 are used during the isolation and characterisation of the genes of the invention.

Sequence 1047 BP; 238 A; 323 C; 299 G; 184 T; 3 other;

Query Match 78.0%; Score 773.6; DB 21; Length 1047;

Best Local Similarity 87.1%; Pred. No. 4.4e-182;

Matches 941; Conservative 1; Mismatches 0; Indels 138; Gaps 3;

```
QY 10 CAGGAAGTGGAGGTTCTCACTGCGCCGAGCAGA-GGCCCTACACCCACGAGGATGGGG 68
DB 1 CAGGAAGTGGAGGTTCTCACTGCGCCGAGCAGAGNGGCCCTACACCCACGAGGATGGGG 60
QY 69 CTCCTCCGCGCTGTTCTGCTGGCGGTGCTGGTGCAGAGGTTCTCCAAAGGACGCGGAG 128
DB 61 CTCCTCCGCGCTGTTCTGCTGGCGGTGCTGGTGCAGAGGTTCTCCAAAGGACGCGGAG 120
QY 129 GAAGAAATTACCCCTGCTGCTCCATTGCTCAAGTCTCGAAGTTTCCCAAGGC 188
DB 121 GAAGAAATTACCCCTGCTGCTCCATTGCTCAAGTCTCGAAGTTTCCCAAGGC 180
QY 189 CGTGGGTGCTCATACCTGCTGTGACCCCGACCCACCCAGGCTTCCATTCCTCC 248
DB 181 CGTGGGTGCTCATACCTGCTGTGACCCCGACCCACCCAGGCTTCCATTCCTCC 240
QY 249 TGTGGAACCAAGAACATCAAGGTGGCCAGAGAGGTGGTGAAGACCCACGAGCGGCTCC 308
DB 241 TGTGGAACCAAGAACATCAAGGTGGCCAGAGAGGTGGTGAAGACCCACGAGCGGCTCC 300
QY 309 TTCAACCTCAAGTCACTCAAGTCCAGTCCAGAGCTGCTCACTTCTGCGGGGG 368
DB 301 TTCAACCTCAAGTCACTCAAGTCCAGTCCAGAGCTGCTCACTTCTGCGGGGG 360
QY 369 TCTTCCACCTCAGGTGCCCATGTGACAGTGGCAGGCTACAGATGACATGGGAGTGTGG 428
DB 361 TCTTCCACCTCAGGTGCCCATGTGACAGTGGCAGGCTACAGATGACATGGGAGTGTGG 420
QY 429 TCAAGCCAGTGTCTGAGTGGCGGGCCAACTTCACTCTGCAAGACAGAGGGGCGGCC 488
DB 421 TCCA-----GACAGAGGGGCGGCC 442
```

```
QY 489 AGGTTGAGATGATCTGCGCAGGCGTCTCTGGGCGAGCCCACTATACCAACAGCCTGATC 548
DB 443 AGGTTGAGATGATCTGCGCAGGCGTCTCTGGGCGAGCCCACTATACCAACAGCCTGATC 502
QY 549 GGGAGGATGGGCGAGGTCCACCTGCGAGCAGAGACCATGCGCAGGCGAGCTGCGCACTTC 608
DB 503 GGGAGGATGGGCGAGGTCCACCTGCGAGCAGAGACCATGCGCAGGCGAGCTGCGCACTTC 562
QY 609 TCCTTCTCTCCGAGCCAGACATCGGACTGCTTCTGCTGCGAGGCTGCAACACAGCCCAAT 668
DB 563 TCCTTCTCTCCGAGCCAGACATCGGACTGCTTCTGCTGCGAGGCTGCAACACAGCCCAAT 622
QY 669 GTCCAGCAGAGCCCTCTCAGTGTGGCC----- 697
DB 623 GTCCAGCAGAGCCCTCTCAGTGTGGTCCGCCAGGAGGTTSCCCAGGGGCACCCACCATC 682
QY 698 ----- 697
DB 683 GTGCTGTTGGCAGCCTTGCTCCACTGCGGGCAATCACCTCCAGGATGTGGGCTGGACC 742
QY 698 -----CCAGGTGGTGACCAAGATGAGGACTGCGAGGTTCCCTGCGAGAGCCCC 749
DB 743 CACGTGGGCCCGAGGTGGTGACCAAGATGAGGACTGCGAGGTTCCCTGCGAGAGCCCC 802
QY 750 ATCTTCTGCTTCCGCTCTTACAGGAGCACCCCGCTCTGAGTGAAGAGAGTGTGGGGGG 809
DB 803 ATCTTCTGCTTCCGCTCTTACAGGAGCACCCCGCTCTGAGTGAAGAGAGTGTGGGGGG 862
QY 810 TTCAAGATAGGAATGGGAGGTCTCAGAGACGCAAGCAGAGCAGGCTGTAGATGAACCG 869
DB 863 TTCAAGATAGGAATGGGAGGTCTCAGAGACGCAAGCAGAGCAGGCTGTAGATGAACCG 922
QY 870 TCCAGAGAGCCAAAGCAGCAGGAGCTCAGGCGCATCAGCGCTGCACTGTTCTATTTGG 929
DB 923 TCCAGAGAGCCAAAGCAGCAGGAGCTCAGGCGCATCAGCGCTGCACTGTTCTATTTGG 982
QY 930 AGTTTCATGCAAAATGAGTGTGTTTAGTGTCTTTCCTGCGCAGGAGGAGGAGGAGGAG 989
DB 983 AGTTTCATGCAAAATGAGTGTGTTTAGTGTCTTTCCTGCGCAGGAGGAGGAGGAGGAG 1042

RESULT 16
ABT16794
ID ABT16794 standard; DNA; 1047 BP.
XX AC ABT16794;
XX XX
DT 03-APR-2003 (first entry)
XX XX
DE Human secreted protein gene sequence - SEQ ID No 43.
XX XX
KW Human; gene; ds; protein therapy; immediate hypersensitivity disease;
allergic disorder; asthmatic disorder; gene therapy; secreted protein;
hay fever; allergic conjunctivitis; allergic rhinitis;
binding partner identification; chromosome identification;
radiation hybrid mapping; long-range restriction mapping.
XX OS Homo sapiens.
XX XX
PN WO200277188-A2.
XX XX
PD 03-OCT-2002.
XX XX
PF 26-MAR-2002; 2002WO-US09239.
XX XX
PR 27-MAR-2001; 2001US-278650P.
PR 12-SEP-2001; 2001US-0950082.
PR 12-SEP-2001; 2001US-0950083.
XX XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX PI Rosen CA, Ruben SM;
XX XX
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DR WPI; 2003-175010/17.
XX
PT Use of human secreted proteins and nucleic acids for preparing a
PT diagnostic or pharmaceutical composition for diagnosing or treating
PT allergic or asthmatic disorders, e.g. asthma, hay fever, or allergic
PT conjunctivitis or rhinitis
XX
PS Claim 7; Page 586; 823pp; English.
XX
CC The invention comprises the amino acid and coding sequences of human
CC secreted proteins. The DNA and protein sequences of the invention are
CC useful for the diagnosis and treatment of allergic disorders, asthmatic
CC disorders and immediate hypersensitivity diseases (e.g. hay fever,
CC allergic conjunctivitis and allergic rhinitis). The proteins of the
CC invention are also useful for identifying a binding partner. The nucleic
CC acids of the invention are also useful for chromosome identification,
CC radiation hybrid mapping or long-range restriction mapping. The present
CC DNA sequence encodes a human secreted protein of the invention.
XX
SQ Sequence 1047 BP; 238 A; 323 C; 299 G; 184 T; 3 other;

Query Match 78.0%; Score 773.6; DB 25; Length 1047;
Best Local Similarity 87.1%; Pred. No. 4.4e-182;
Matches 941; Conservative 1; Mismatches 0; Indels 138; Gaps 3;

QY 10 CAGGAAGTGGAGGTTCTCACTGCCCGAGCAGA-GGCCCTACACCCACCGAGGCAATGGG 68
DB 1 CAGGAAGTGGAGGTTCTCACTGCCCGAGCAGA-GGCCCTACACCCACCGAGGCAATGGG 60
QY 69 CTCCTGGGCTTTCTGCTGGCGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 128
DB 61 CTCCTGGGCTTTCTGCTGGCGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 120
QY 129 GAAGAAATACCCCTGTGCTCTCCATTGCTCAAGTCTCAAGTCTCAAGTCTCAAGTCTCA 188
DB 121 GAAGAAATACCCCTGTGCTCTCCATTGCTCAAGTCTCAAGTCTCAAGTCTCAAGTCTCA 180
QY 189 CGCTGGGTGCTCATACCTGCTGTGTCACCCAGCCACACCCAGCCACCTATTCCTTCCTTC 248
DB 181 CGCTGGGTGCTCATACCTGCTGTGTCACCCAGCCACACCCAGCCACCTATTCCTTCCTTC 240
QY 249 TGTGGAACCAAGAACATCAAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 308
DB 241 TGTGGAACCAAGAACATCAAGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGGTGG 300
QY 309 TTCAACCTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCA 368
DB 301 TTCAACCTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCAAGTCA 360
QY 369 TCCTCCACTCAGGTGCCCATGTGACAGTGCAGGCTACAGATGACCTGGGAGCTGTGG 428
DB 361 TCCTCCACTCAGGTGCCCATGTGACAGTGCAGGCTACAGATGACCTGGGAGCTGTGG 420
QY 429 TCCAGGCCAGTCTTGAGCTGGGGCCCACTTCACTCTGCAGACAGAGGGGACAGGCCCC 488
DB 421 TCCA-----GACAGAGGGGACAGGCCCC 442
QY 489 AGGGTGGAGATGATCTGCCAGGCGTCTCGGGCAGCCCACTTATCAACCAAGCCTGATC 548
DB 443 AGGGTGGAGATGATCTGCCAGGCGTCTCGGGCAGCCCACTTATCAACCAAGCCTGATC 502
QY 549 GGAAGAGTGGGAGTCCACTGTCAGCAGAGACCATGTCACAGGAGCTGCCAATTC 608
DB 503 GGAAGAGTGGGAGTCCACTGTCAGCAGAGACCATGTCACAGGAGCTGCCAATTC 562
QY 609 TCCTTCCTGCCAGCCAGACATCGACTGTTCTGTTGTCAGGCTGCAACAGCCCAAT 668
DB 563 TCCTTCCTGCCAGCCAGACATCGACTGTTCTGTTGTCAGGCTGCAACAGCCCAAT 622
QY 669 GTCCAGCAGAGCCCTCTCAGTGGTGGC----- 697
DB 623 GTCCAGCAGAGCCCTCTCAGTGGTGGC----- 697
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QY 698 ----- 697
DB 683 GTGCTGTTGSCAGGCTTGCTCCACTCGCGGCATCACTCCAGGATGCTGGGCTGGACC 742
QY 698 -----CCAGGTGGTGACCAAGATGGAGGACTGGCAGGGTCCCTCGAGAGCCCC 749
DB 743 CACGTGGGCCCCAGGTGGTGACCAAGATGGAGGACTGGCAGGGTCCCTCGAGAGCCCC 802
QY 750 ATCCTTGCTTGCCGCTCTACAGGACACCCGCTCTGAGTGGAGAGAGTTTGGGGGG 809
DB 803 ATCCTTGCTTGCCGCTCTACAGGACACCCGCTCTGAGTGGAGAGAGTTTGGGGGG 862
QY 810 TTCAGGATAGGAATGGGGAGGTGACAGGACGCAAGCAGAGCCATGTAGAAATCAACCG 869
DB 863 TTCAGGATAGGAATGGGGAGGTGACAGGACGCAAGCAGAGCCATGTAGAAATCAACCG 922
QY 870 TCCAGAGCCCAAGCAGCAGAGGACTGTCAGGCGCATCAGCGTGCATCTGTCGTATTGG 929
DB 923 TCCAGAGCCCAAGCAGCAGAGGACTGTCAGGCGCATCAGCGTGCATCTGTCGTATTGG 982
QY 930 AGTTTCATGCAAAATGAGTGTCTTTTAGCTGCTTGCACAAAAAAGGAGAGAGAGAG 989
DB 983 AGTTTCATGCAAAATGAGTGTCTTTTAGCTGCTTGCACAAAAAAGGAGAGAGAGAG 1042

RESULT 17
ABZ66993
ID ABZ66993 standard; cDNA; 1047 BP.
XX
AC ABZ66993;
XX
DT 26-MAR-2003 (first entry)
XX
DE Human secreted protein encoding cDNA SEQ ID NO 113.
XX
KW Human; secreted protein; nootropic; neuroprotective; cytostatic;
KW viricide; dermatological; immunosuppressive; antiinflammatory; anti-HIV;
KW vulnary; antibacterial; antiparkinsonian; antiskinning; antianemic;
KW antiarthritic; cancer; antirheumatic; hepatotropic; cerebroprotective;
KW antiinflammatory; antiallergic; antidiabetic; antituber; anticonvulsant;
KW antifungal; antiparasitic; cardiac; immune disorder; infection; vaccine;
KW cardiovascular disorder; neurological disease; nephrotropic;
KW gene therapy; gene; ds.
XX
OS Homo sapiens.
XX
PN WO200277186-A2.
XX
PD 03-OCT-2002.
XX
PF 26-MAR-2002; 2002WO-US09188.
XX
PR 27-MAR-2001; 2001US-278650P.
XX
PR 12-SEP-2001; 2001US-0950082.
XX
PR 12-SEP-2001; 2001US-0950083.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Rosen CA, Ruben SM;
XX
XX WPI; 2003-040583/03.
XX
DR P-PSDB; ABP99572.
XX
PT New human secreted proteins encoded by genes contained in cDNA clones
PT (e.g. HGCA19), useful for preventing, treating or diagnosing e.g.
PT AIDS, multiple sclerosis, herpes virus, leukemia, tick-borne
PT encephalitis or West Nile fever
XX
PS Claim 7; Page 1235-1236; 2423pp; English.
XX
CC The invention relates to novel human genes (ABZ66891-ABZ68209) and the
CC encoded secreted proteins (ABP99470-ABP99872) useful for preventing,
CC treating or ameliorating medical conditions e.g. by protein or gene
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CC therapy. The genes are isolated from a range of human tissues disclosed
CC in the specification. The nucleic acids, proteins, antibodies and
CC (ant)agonists are useful in the diagnosis, treatment and prevention of:
CC (a) cancer, e.g. breast and ovarian cancer and other cancers of the
CC adrenal gland, bone, bone marrow, breast, gastrointestinal tract, liver,
CC lung or urogenital; (b) immune disorders e.g. Addison's disease,
CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
CC arthritis and ulcerative colitis; (c) cardiovascular diseases such as
CC myocardial ischaemias; (d) wound healing; (e) neurological diseases e.g.
CC cerebral anoxia and epilepsy; and (f) infectious diseases such as viral,
CC bacterial, fungal and parasitic infections.

XX SQ Sequence 1047 BP; 238 A; 323 C; 299 G; 184 T; 3 other;

Query Match 78.0%; Score 773.6; DB 25; Length 1047;
Best Local Similarity 87.1%; Pred. No. 4.4e-182;
Matches 941; Conservative 1; Mismatches 0; Indels 138; Gaps 3;

```
QY 10 CAGGAATAGAGGTTCTCTACGCGGAGCAGA-GGCCCTACACCCACCGAGGATGGG 68
D 1 CAGGAATAGAGGTTCTCTACGCGGAGCAGA-GGCCCTACACCCACCGAGGATGGG 60
QY 69 CTCCTGGGCTGTTCTGCTTGGCGTCTGCTGCGAGAGCTTCTCCAGGACAGGGAG 128
D 61 CTCCTGGGCTGTTCTGCTTGGCGTCTGCTGCGAGAGCTTCTCCAGGACAGGGAG 120
QY 129 GAAGAAATACCCCTGTGGTCTCATGTGCTCAAGTCTCTGGAAGTTTCCCAAGGC 188
D 121 GAAGAAATACCCCTGTGGTCTCATGTGCTCAAGTCTCTGGAAGTTTCCCAAGGC 180
QY 189 CGCTGGGTGTCATAAAGTCTGTGACCCGACCCAGCCGACCTATTCCTTC 248
D 181 CGCTGGGTGTCATAAAGTCTGTGACCCGACCCAGCCGACCTATTCCTTC 240
QY 249 TGTGGACCAAGACATCAGGTGGCCAGAAAGTGTGAAGCCACGAGCGGCTCC 308
D 241 TGTGGACCAAGACATCAGGTGGCCAGAAAGTGTGAAGCCACGAGCGGCTCC 300
QY 309 TTCAAGCTCAACCTCACTCAAGTCCAGTCCAGACCTGCTCACTTCTGCCGGCG 368
D 301 TTCAAGCTCAACCTCACTCAAGTCCAGTCCAGACCTGCTCACTTCTGCCGGCG 360
QY 369 TCTTCACTCAAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGT 428
D 361 TCTTCACTCAAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGTCCAGT 420
QY 429 TCCAAGCCAGTGTGAGCTGCGGCGCACTTCTCACTGTCAGGACAGGGGCGGCG 488
D 421 TCCAAGCCAGTGTGAGCTGCGGCGCACTTCTCACTGTCAGGACAGGGGCGGCG 442
QY 489 AGGTGGAGATGATCTGCCAGGCGTCTCGGAGCGCCCTATCACTCAACACCTGATC 548
D 443 AGGTGGAGATGATCTGCCAGGCGTCTCGGAGCGCCCTATCACTCAACACCTGATC 502
QY 549 GGAAGAGTGGGAGGTCCACCTGTCAGCAGACCAATGCCAGGAGCGCTGCCACTTC 608
D 503 GGAAGAGTGGGAGGTCCACCTGTCAGCAGACCAATGCCAGGAGCGCTGCCACTTC 562
QY 609 TCTTCTCGGAGCAGACATCGGAGTGTCTGTCGTCAGGCTGCAACACGCAAT 668
D 563 TCTTCTCGGAGCAGACATCGGAGTGTCTGTCGTCAGGCTGCAACACGCAAT 622
QY 669 GTCCAGCAGCGCCCTCAAGTGGTGGC----- 697
D 623 GTCCAGCAGCGCCCTCAAGTGGTGGC----- 682
QY 698 ----- 697
D 683 GTGCTGTTGGAGCCTTGGCTCCACTGCGGCGCATCACCTCCAGGATGTGGGCTGGAC 742
QY 698 -----CCAGTGGTACACAGATGAGAGCTGGCAGGTTCCCTGGAGAGCCCC 749
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Db 743 CACGTGGGCCAGGTGTGTGACCAAGATGGAGGACTGGCAGGTCCTCTGGAGAGCCCC 802
QY 750 ATCCCTTGCCCTTCCGCTCTACAGGACACCCCGCTGTGAGTGAAGAGGAGTTTGGGGG 809
D 803 ATCCCTTGCCCTTCCGCTCTACAGGACACCCCGCTGTGAGTGAAGAGGAGTTTGGGGG 862
QY 810 TTCAGGATAGGGAATGGGAGGTTCAGAGGACGCAAGACGACGACCATGTAGAATGAACCG 869
D 863 TTCAGGATAGGGAATGGGAGGTTCAGAGGACGCAAGACGACGACCATGTAGAATGAACYG 922
QY 870 TCAGAGAGCCAGACGCGGAGGACTGCGAGGCGCATCAGCGTGCACCTGTTCGTATTTGG 929
D 923 TCAGAGAGCCAGACGCGGAGGACTGCGAGGCGCATCAGCGTGCACCTGTTCGTATTTGG 982
QY 930 AGTTTCATGCAAAATGAGTGTGTTTGTAGTCTCTTCCACAAAAAAGGAGGAGGAGG 989
D 983 AGTTTCATGCAAAATGAGTGTGTTTGTAGTCTCTTCCACAAAAAAGGAGGAGGAGG 1042
```

RESULT 18

ABZ36822

ID ABZ36822 standard; cDNA; 502 BP.

AC ABZ36822;

XX 21-FEB-2003 (first entry)

XX Human GENSET coding sequence SEQ ID 695.

XX Cytostatic; antiinflammatory; nootropic; neuroprotective; cardiac;
KW gastrointestinal; gene therapy; GENSET; heavy metal toxicity; cancer;
KW inflammatory disease; immune disorder; neuromuscular; toxicity;
KW central nervous system; cardiovascular; gastrointestinal; gene; ss.

OS Homo sapiens.

PN W0200283898-A1.

XX 24-OCT-2002.

XX 18-APR-2001; 2001WO-IB00914.

XX 18-APR-2001; 2001WO-IB00914.

XX (GENSET) GENSET.

PI Bejanin S, Tanaka H, Dumas Milne Edwards J, Jobert S, Giordano J;
XX WPI; 2003-075548/07.

DR New GENSET polynucleotides and polypeptides, useful for treating heavy
PT metal toxicity, cancer, inflammatory diseases, immune disorders, and
PT the neuromuscular, CNS, cardiovascular or gastrointestinal effects of
PT the toxicity -

PS Claim 12; Page 656; 735pp; English.

XX The present invention relates to novel GENSET polynucleotides
CC (ABZ36404-ABZ36911) encoding polypeptides (ABP75963-ABP76368). The
CC polynucleotides and polypeptides are useful in screening and diagnostic
CC assays for abnormal GENSET expression and/or biological activity. They
CC are also useful for screening of compounds for treating or preventing
CC GENSET-related disorders, such as heavy metal toxicity, cancer,
CC inflammatory diseases, immune disorders, and the neuromuscular, central
CC nervous system (CNS), cardiovascular or gastrointestinal effects of the
CC toxicity. The polynucleotides are useful for constructing or expanding
CC chromosome maps.

SQ Sequence 502 BP; 105 A; 169 C; 135 G; 92 T; 1 other;

Query Match 50.6%; Score 501.6; DB 25; Length 502;

Best Local Similarity 99.8%; Pred. No. 1.1e-114;

Matches 501; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

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QY 40 AGAGGCCCTACACCAAGGAGGATGGGGTCCCTGGGCTGTCTCTTGGCGCTGCTGG 99
Db 1 AGAGGCCCTACACCAAGGAGGATGGGGTCCCTGGGCTGTCTCTTGGCGCTGCTGG 60
QY 100 CTGCCAGCAGCTTCTCCAAAGGACGGGAGGAGAAATTACCCCTGGTGTCTCCATTGCTT 159
Db 61 CTGCCAGCAGCTTCTCCAAAGGACGGGAGGAGAAATTACCCCTGGTGTCTCCATTGCTT 120
QY 160 ACAAGTCTGTGAAGTTTTCACAAAGGCGCTGGTGTCTATAACCTGTCTGTGCACCC 219
Db 121 ACAAGTCTGTGAAGTTTTCACAAAGGCGCTGGTGTCTATAACCTGTCTGTGCACCC 180
QY 220 AGCCACCAAGCCCATCACTATTCCTCTGTGGAACCAAGAACATCAAGGTGGCCAGA 279
Db 181 AGCCACCAAGCCCATCACTATTCCTCTGTGGAACCAAGAACATCAAGGTGGCCAGA 240
QY 280 AGTGTGTGAAGACCAAGGCGGCTCTCTCAACCTCAACGTCACACTCAAGTCCAGTC 339
Db 241 AGTGTGTGAAGACCAAGGCGGCTCTCTCAACCTCAACGTCACACTCAAGTCCAGTC 300
QY 340 CAGACTGTCTACCTACTTCTCCGGGGTCTCTCAACCTCAGGTGCCATGTGGACAGTG 399
Db 301 CAGACTGTCTACCTACTTCTCCGGGGTCTCTCAACCTCAGGTGCCATGTGGACAGTG 360
QY 400 CCAGGCTACAGTACACTGGGAGCTGTGTCCAAGCAGTGTCTGAGTGGCGGCAACT 459
Db 361 CCAGGCTACAGTACACTGGGAGCTGTGTCCAAGCAGTGTCTGAGTGGCGGCAACT 420
QY 460 TCACCTGTGAGACAGAGGGGAGGCGGCTCTCAACCTCAACGTCAGATGATCGCAGGGGCTCTCGG 519
Db 421 TCACCTGTGAGACAGAGGGGAGGCGGCTCTCAACCTCAACGTCAGATGATCGCAGGGGCTCTCGG 480
QY 520 GCAGGCCCTATCAACCAAG 541
Db 481 GCAGGCCCTATCAACCAAG 502

RESULT 19
AAS34076
ID AAS34076 standard; cDNA; 724 BP.
AC AAS34076;
DT 17-DEC-2001 (first entry)
DE Human cDNA encoding a novel foetal antigen, SEQ ID No 600.
KW Human; foetal tissue antigen; ss; antiinflammatory; neuroprotective;
KW immunomodulator; cardiovascular; cytostatic; nephrothropic;
KW cardiovascular; autoimmune disease; rheumatoid arthritis;
KW hyperproliferative disorder; breast neoplasm; cancer;
KW cardiovascular disorder; cardiac arrest; cerebrovascular disorder;
KW cerebral ischaemia; angiogenesis; nervous system disorder;
KW Alzheimer's disease; infection; ocular disorder; corneal infection;
KW wound healing; epithelial cell proliferation; food additive.
XX OS Homo sapiens.
XX
XX WO200155312-A2.
XX
XX 02-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01321.
XX
XX 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189874.
XX 17-MAR-2000; 2000US-0190076.
XX 18-APR-2000; 2000US-0198123.
XX 19-MAY-2000; 2000US-0205515.
XX
XX 07-JUN-2000; 2000US-0209467.
XX 28-JUN-2000; 2000US-0214886.
XX 30-JUN-2000; 2000US-0215135.
XX 07-JUL-2000; 2000US-0216647.
XX 07-JUL-2000; 2000US-0216880.
XX 11-JUL-2000; 2000US-0217487.
XX 11-JUL-2000; 2000US-0217496.
XX 14-JUL-2000; 2000US-0218290.
XX 26-JUL-2000; 2000US-0220963.
XX 26-JUL-2000; 2000US-0220964.
XX 14-AUG-2000; 2000US-0224518.
XX 14-AUG-2000; 2000US-0224519.
XX 14-AUG-2000; 2000US-0225213.
XX 14-AUG-2000; 2000US-0225214.
XX 14-AUG-2000; 2000US-0225266.
XX 14-AUG-2000; 2000US-0225267.
XX 14-AUG-2000; 2000US-0225268.
XX 14-AUG-2000; 2000US-0225270.
XX 14-AUG-2000; 2000US-0225447.
XX 14-AUG-2000; 2000US-0225757.
XX 14-AUG-2000; 2000US-0225758.
XX 14-AUG-2000; 2000US-0225759.
XX 18-AUG-2000; 2000US-0226279.
XX 22-AUG-2000; 2000US-0226681.
XX 22-AUG-2000; 2000US-0226868.
XX 22-AUG-2000; 2000US-0227182.
XX 23-AUG-2000; 2000US-0227009.
XX 30-AUG-2000; 2000US-0228924.
XX 01-SEP-2000; 2000US-0229287.
XX 01-SEP-2000; 2000US-0229343.
XX 01-SEP-2000; 2000US-0229344.
XX 01-SEP-2000; 2000US-0229345.
XX 05-SEP-2000; 2000US-0229509.
XX 05-SEP-2000; 2000US-0229513.
XX 06-SEP-2000; 2000US-0230437.
XX 06-SEP-2000; 2000US-0230438.
XX 08-SEP-2000; 2000US-0231242.
XX 08-SEP-2000; 2000US-0231243.
XX 08-SEP-2000; 2000US-0231244.
XX 08-SEP-2000; 2000US-0231413.
XX 08-SEP-2000; 2000US-0231414.
XX 08-SEP-2000; 2000US-0232080.
XX 12-SEP-2000; 2000US-0231968.
XX 14-SEP-2000; 2000US-0232397.
XX 14-SEP-2000; 2000US-0232398.
XX 14-SEP-2000; 2000US-0232399.
XX 14-SEP-2000; 2000US-0232400.
XX 14-SEP-2000; 2000US-0232401.
XX 14-SEP-2000; 2000US-0233063.
XX 14-SEP-2000; 2000US-0233064.
XX 21-SEP-2000; 2000US-0233065.
XX 21-SEP-2000; 2000US-0234223.
XX 21-SEP-2000; 2000US-0234274.
XX 25-SEP-2000; 2000US-0234997.
XX 25-SEP-2000; 2000US-0234998.
XX 26-SEP-2000; 2000US-0235484.
XX 27-SEP-2000; 2000US-0235834.
XX 27-SEP-2000; 2000US-0235836.
XX 29-SEP-2000; 2000US-0236327.
XX 29-SEP-2000; 2000US-0236367.
XX 29-SEP-2000; 2000US-0236368.
XX 29-SEP-2000; 2000US-0236369.
XX 29-SEP-2000; 2000US-0236370.
XX 02-OCT-2000; 2000US-0236802.
XX 02-OCT-2000; 2000US-0237037.
XX 02-OCT-2000; 2000US-0237038.
XX 02-OCT-2000; 2000US-0237039.
XX 02-OCT-2000; 2000US-0237040.
XX 13-OCT-2000; 2000US-0239935.
XX 13-OCT-2000; 2000US-0239937.
XX 20-OCT-2000; 2000US-0240960.
XX 20-OCT-2000; 2000US-0241221.
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PR 20-OCT-2000; 2000US-0241785.
 PR 20-OCT-2000; 2000US-0241786.
 PR 20-OCT-2000; 2000US-0241787.
 PR 20-OCT-2000; 2000US-0241788.
 PR 20-OCT-2000; 2000US-0241808.
 PR 20-OCT-2000; 2000US-0241809.
 PR 20-OCT-2000; 2000US-0241826.
 PR 01-NOV-2000; 2000US-0244617.
 PR 08-NOV-2000; 2000US-0246474.
 PR 08-NOV-2000; 2000US-0246475.
 PR 08-NOV-2000; 2000US-0246476.
 PR 08-NOV-2000; 2000US-0246477.
 PR 08-NOV-2000; 2000US-0246478.
 PR 08-NOV-2000; 2000US-0246523.
 PR 08-NOV-2000; 2000US-0246524.
 PR 08-NOV-2000; 2000US-0246525.
 PR 08-NOV-2000; 2000US-0246526.
 PR 08-NOV-2000; 2000US-0246527.
 PR 08-NOV-2000; 2000US-0246528.
 PR 08-NOV-2000; 2000US-0246532.
 PR 08-NOV-2000; 2000US-0246609.
 PR 08-NOV-2000; 2000US-0246610.
 PR 08-NOV-2000; 2000US-0246611.
 PR 08-NOV-2000; 2000US-0246613.
 PR 17-NOV-2000; 2000US-0249207.
 PR 17-NOV-2000; 2000US-0249208.
 PR 17-NOV-2000; 2000US-0249209.
 PR 17-NOV-2000; 2000US-0249210.
 PR 17-NOV-2000; 2000US-0249211.
 PR 17-NOV-2000; 2000US-0249212.
 PR 17-NOV-2000; 2000US-0249213.
 PR 17-NOV-2000; 2000US-0249214.
 PR 17-NOV-2000; 2000US-0249215.
 PR 17-NOV-2000; 2000US-0249216.
 PR 17-NOV-2000; 2000US-0249217.
 PR 17-NOV-2000; 2000US-0249218.
 PR 17-NOV-2000; 2000US-0249244.
 PR 17-NOV-2000; 2000US-0249245.
 PR 17-NOV-2000; 2000US-0249264.
 PR 17-NOV-2000; 2000US-0249265.
 PR 17-NOV-2000; 2000US-0249297.
 PR 17-NOV-2000; 2000US-0249299.
 PR 01-DEC-2000; 2000US-0250160.
 PR 05-DEC-2000; 2000US-0250391.
 PR 05-DEC-2000; 2000US-0251030.
 PR 05-DEC-2000; 2000US-0251988.
 PR 05-DEC-2000; 2000US-0256719.
 PR 08-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
 PR 08-DEC-2000; 2000US-0251868.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 08-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX
 XX Rosen CA, Barash SC, Ruben SM;
 XX WPI; 2001-488782/53.
 XX P-PSDB; AAU21256.
 XX
 XX New polynucleotides and polypeptides for diagnosing, treating,
 XX preventing or prognosing e.g. diseases or disorders of the nervous,
 XX musculoskeletal, excretory, gastrointestinal, reproductive, and
 XX respiratory systems -
 XX
 XX Claim 1; SEQ ID No 600; 642pp; English.
 XX
 XX The invention relates to novel nucleic acids encoding novel human foetal
 XX antigens. The nucleic acids and proteins are used to prevent, treat (e.g.
 XX by gene therapy) or ameliorate a medical condition in e.g. humans, mice,

CC rabbits, goats, horses, cats, dogs, chickens or sheep. They
 CC are also used in diagnosing a pathological condition or susceptibility
 CC to a pathological condition. The antibodies to the antigens can also
 CC be used in alleviating symptoms associated with the disorders and in
 CC diagnostic immunoassays e.g. radioimmunoassays or enzyme linked
 CC immunosorbent assays (ELISA). Disorders which are diagnosed or treated
 CC include autoimmune diseases e.g. rheumatoid arthritis,
 CC hyperproliferative disorders e.g. neoplasms of the breast or liver,
 CC cardiovascular disorders e.g. cardiac arrest, cerebrovascular disorders
 CC e.g. cerebral ischaemia, angiogenesis, nervous system disorders e.g.
 CC Alzheimer's disease, infections caused by bacteria, viruses and fungi
 CC and ocular disorders e.g. corneal infection. The polypeptides can also
 CC be used to aid wound healing and epithelial cell proliferation, to
 CC prevent skin aging due to sunburn, to maintain organs before
 CC transplantation, for supporting cell culture of primary tissues, to
 CC regenerate tissues and in chemotaxis. The polypeptides can also be used
 CC as a food additive or preservative to increase or decrease storage
 CC capabilities, fat content, lipid, protein, carbohydrate, vitamins,
 CC minerals, cofactors and other nutritional components. Numerous
 CC examples of diseases and disorders treated by the nucleic acids and
 CC proteins are given in the specification. The present sequence

Query Match 50.6%; Score 501.6; DB 22; Length 724;
 Best Local Similarity 98.0%; Pred. No. 1.2e-114;

Matches 545; Conservative 4; Mismatches 3; Indels 4; Gaps 4;

QY	131	AGAAATTACCCCTGTGCTCTCCATTGCTTACAAAGTCTTGGAAAGTTTCCCAAGGCGG	190
DB	161	AGAAATTACCCCTGTGCTCTCCATTGCTTACAAAGTCTTGGAAAGTTTCCCAAGGCGG	220
QY	191	CTGGGTGCTCATAAOCCTGCTGTGACCCAGCCACCCAGGCTTACCTATTCCTCTG	250
DB	221	CTGGGTGCTCATAAOCCTGCTGTGACCCAGCCACCCAGGCTTACCTATTCCTCTG	280
QY	251	TGGAAACCAAGACATCAAGTGGCCAAAGAGTGGTGAAGACCCAGAGCGGCTCTCT	310
DB	281	TGGAAACCAAGACATCAAGTGGCCAAAGAGTGGTGAAGACCCAGAGCGGCTCTCT	340
QY	311	CAACCTCAACGTCACACTCAAGTCCAGTCCAGACCTGCTCAGTACTTCTGCGGCGTC	370
DB	341	CAACCTCAACGTCACACTCAAGTCCAGTCCAGACCTGCTCAGTACTTCTGCGGCGTC	400
QY	371	CTCCACCTCAGTGGCCCATGTGGACAGTGCAGGCTACAGATGCCATGGGAGCTGTGTC	430
DB	401	CTCCACCTCAGTGGCCCATGTGGACAGTGCAGGCTACAGATGCCATGGGAGCTGTGTC	460
QY	431	CAAGCCAGTGTGAGCTGCGGGCCAACTTCACTCTGCAAGGACAGAGGGGAGGCCCCAG	490
DB	461	CAAGCCAGTGTGAGCTGCGGGCCAACTTCACTCTGCAAGGACAGAGGGGAGGCCCCAG	519
QY	491	GGTGGAGATGATCTGCCAGGCGTCTCGGGGAGCCCACTATCACCACAGCCTGATCGG	550
DB	520	GGTGGAGATGATCTGCCAGGCGTCTCGGGGAGCCCACTATCACCACAGCCTGATCGG	578
QY	551	GAAGGATGGGAGGTTCCACCTGTCAGCAGAGACCATGSCCAGAGGAGCTTCCAACTTCTC	610
DB	579	GAAGGATGGGAGGTTCCACCTGTCAGCAGAGACCATGSCCAGAGGAGCTTCCAACTTCTC	637
QY	611	CTTCTGCGGAGCCAGACATCGGAGTGTTC-TGGTGCAGGCTGCAAAACAGCGCAATG	669
DB	638	CTTCTGCGGAGCCAGACATCGGAGTGTTC-TGGTGCAGGCTGCAAAACAGCGCAATG	697
QY	670	TCCAGCACAGCGCCCT 685	
DB	698	TCCAGCACAGCGCCCT 713	

RESULT 20
 ABA20618

ID ABA20618 standard; DNA; 32220 BP.

XX AC ABA20618;
 XX

DT 23-JAN-2002 (first entry)
XX Human nervous system related polynucleotide SEQ ID NO 12949.
DE XX
XX Human; nontropic; neuroprotective; cytostatic; dermatological; virucide;
KW immunosuppressive; antiinflammatory; anti-HIV; antibacterial; vulnerary;
KW antiparkinsonian; antiskelting; antianemic; antiarthritic; cancer;
KW antirheumatic; hepatotropic; cerebroprotective; antiinflammatory;
KW antiallergic; antidiabetic; antifungal; anticonvulsant; antifungal;
KW antiparasitic; cardiac; immune disorder; cardiovascular disorder;
KW neurological disease; infection; nephrotropic; gene therapy; vaccine; ds.
XX OS
XX Homo sapiens.
XX WO200159063-A2.
XX 16-AUG-2001.
XX
XX 17-JAN-2001; 2001WO-US01334.
XX
XX 31-JAN-2000; 2000US-0179065.
XX 04-FEB-2000; 2000US-0180628.
XX 24-FEB-2000; 2000US-0184664.
XX 02-MAR-2000; 2000US-0186350.
XX 16-MAR-2000; 2000US-0189874.
XX 17-MAR-2000; 2000US-0190076.
XX 18-APR-2000; 2000US-0198123.
XX 19-MAY-2000; 2000US-0205515.
XX 07-JUN-2000; 2000US-0209467.
XX 28-JUN-2000; 2000US-0214886.
XX 30-JUN-2000; 2000US-0215135.
XX 07-JUL-2000; 2000US-0215647.
XX 07-JUL-2000; 2000US-0216680.
XX 11-JUL-2000; 2000US-0217487.
XX 11-JUL-2000; 2000US-0217496.
XX 14-JUL-2000; 2000US-0218290.
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XX 14-AUG-2000; 2000US-0225266.
XX 14-AUG-2000; 2000US-0225267.
XX 14-AUG-2000; 2000US-0225268.
XX 14-AUG-2000; 2000US-0225270.
XX 14-AUG-2000; 2000US-0225271.
XX 14-AUG-2000; 2000US-0225275.
XX 14-AUG-2000; 2000US-0225757.
XX 14-AUG-2000; 2000US-0225758.
XX 14-AUG-2000; 2000US-0225759.
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XX 22-AUG-2000; 2000US-0226681.
XX 22-AUG-2000; 2000US-0226686.
XX 22-AUG-2000; 2000US-0227182.
XX 23-AUG-2000; 2000US-0227009.
XX 30-AUG-2000; 2000US-0228924.
XX 01-SEP-2000; 2000US-0229287.
XX 01-SEP-2000; 2000US-0229343.
XX 01-SEP-2000; 2000US-0229344.
XX 01-SEP-2000; 2000US-0229345.
XX 05-SEP-2000; 2000US-0229509.
XX 05-SEP-2000; 2000US-0229513.
XX 06-SEP-2000; 2000US-0230437.
XX 06-SEP-2000; 2000US-0230438.
XX 08-SEP-2000; 2000US-0231242.
XX 08-SEP-2000; 2000US-0231243.
XX 08-SEP-2000; 2000US-0231244.
XX 08-SEP-2000; 2000US-0231413.
XX 08-SEP-2000; 2000US-0231414.
XX 08-SEP-2000; 2000US-0232080.
XX 08-SEP-2000; 2000US-0232081.
XX 12-SEP-2000; 2000US-0231968.
XX 14-SEP-2000; 2000US-0232397.
PR 14-SEP-2000; 2000US-0232398.
PR 14-SEP-2000; 2000US-0232399.
PR 14-SEP-2000; 2000US-0232400.
PR 14-SEP-2000; 2000US-0232401.
PR 14-SEP-2000; 2000US-0233063.
PR 14-SEP-2000; 2000US-0233064.
PR 14-SEP-2000; 2000US-0233065.
PR 21-SEP-2000; 2000US-0234223.
PR 21-SEP-2000; 2000US-0234224.
PR 25-SEP-2000; 2000US-0234997.
PR 25-SEP-2000; 2000US-0234998.
PR 26-SEP-2000; 2000US-0234984.
PR 27-SEP-2000; 2000US-0235834.
PR 27-SEP-2000; 2000US-0235836.
PR 29-SEP-2000; 2000US-0236327.
PR 29-SEP-2000; 2000US-0236367.
PR 29-SEP-2000; 2000US-0236368.
PR 29-SEP-2000; 2000US-0236369.
PR 29-SEP-2000; 2000US-0236370.
PR 02-OCT-2000; 2000US-0236802.
PR 02-OCT-2000; 2000US-0237037.
PR 02-OCT-2000; 2000US-0237038.
PR 02-OCT-2000; 2000US-0237039.
PR 13-OCT-2000; 2000US-0239935.
PR 13-OCT-2000; 2000US-0239937.
PR 20-OCT-2000; 2000US-0240960.
PR 20-OCT-2000; 2000US-0241785.
PR 20-OCT-2000; 2000US-0241786.
PR 20-OCT-2000; 2000US-0241787.
PR 20-OCT-2000; 2000US-0241808.
PR 20-OCT-2000; 2000US-0241809.
PR 20-OCT-2000; 2000US-0241826.
PR 01-NOV-2000; 2000US-0242221.
PR 08-NOV-2000; 2000US-0244617.
PR 08-NOV-2000; 2000US-0246474.
PR 08-NOV-2000; 2000US-0246475.
PR 08-NOV-2000; 2000US-0246476.
PR 08-NOV-2000; 2000US-0246477.
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PR 08-NOV-2000; 2000US-0246525.
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PR 08-NOV-2000; 2000US-0246527.
PR 08-NOV-2000; 2000US-0246528.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246532.
PR 08-NOV-2000; 2000US-0246609.
PR 08-NOV-2000; 2000US-0246610.
PR 08-NOV-2000; 2000US-0246611.
PR 08-NOV-2000; 2000US-0246613.
PR 17-NOV-2000; 2000US-0249207.
PR 17-NOV-2000; 2000US-0249208.
PR 17-NOV-2000; 2000US-0249209.
PR 17-NOV-2000; 2000US-0249210.
PR 17-NOV-2000; 2000US-0249211.
PR 17-NOV-2000; 2000US-0249212.
PR 17-NOV-2000; 2000US-0249213.
PR 17-NOV-2000; 2000US-0249214.
PR 17-NOV-2000; 2000US-0249215.
PR 17-NOV-2000; 2000US-0249216.
PR 17-NOV-2000; 2000US-0249217.
PR 17-NOV-2000; 2000US-0249218.
PR 17-NOV-2000; 2000US-0249244.
PR 17-NOV-2000; 2000US-0249245.
PR 17-NOV-2000; 2000US-0249246.
PR 17-NOV-2000; 2000US-0249265.
PR 17-NOV-2000; 2000US-0249297.
PR 17-NOV-2000; 2000US-0249299.
PR 17-NOV-2000; 2000US-0249300.
PR 01-DEC-2000; 2000US-0250391.
PR 01-DEC-2000; 2000US-0251160.
PR 05-DEC-2000; 2000US-0251030.

PR 05-DEC-2000; 2000US-0251988.
 PR 06-DEC-2000; 2000US-0256719.
 PR 06-DEC-2000; 2000US-0251479.
 PR 08-DEC-2000; 2000US-0251856.
 PR 08-DEC-2000; 2000US-0251868.
 PR 08-DEC-2000; 2000US-0251869.
 PR 08-DEC-2000; 2000US-0251989.
 PR 08-DEC-2000; 2000US-0251990.
 PR 11-DEC-2000; 2000US-0254097.
 PR 05-JAN-2001; 2001US-0259678.
 XX (HUMA-) HUMAN GENOME SCI INC.
 XX Rosen CA, Barash SC, Ruben SM;
 XX WPI; 2001-541565/60.
 XX
 PT Nucleic acids encoding 3224 human nervous system antigen polypeptides,
 PT useful for preventing, diagnosing and/or treating nervous system
 PT cancers and metastases -
 XX
 PS Disclosure; SEQ ID NO 12949; 1701pp + Sequence Listing; English.
 XX
 CC The invention relates to novel genes (ABA11004-ABA21534) and proteins
 CC (AB114678-AB18001) useful for preventing, treating or ameliorating
 CC medical conditions e.g. by protein or gene therapy. The genes are
 CC isolated from a range of human tissues disclosed in the specification.
 CC The nucleic acids, proteins, antibodies and (ant)agonists are useful
 CC in the diagnosis, treatment and prevention of: (a) cancer, e.g. breast
 CC and ovarian cancer and other cancers of the adrenal gland, bone, bone
 CC marrow, breast, gastrointestinal tract, liver, lung, or urogenital;
 CC (b) immune disorders e.g. Addison's disease, allergies, autoimmune
 CC haemolytic anaemia, autoimmune thyroiditis, diabetes mellitus, Crohn's
 CC disease, multiple sclerosis, rheumatoid arthritis and ulcerative
 CC colitis; (c) cardiovascular disorders such as myocardial ischaemias;
 CC (d) wound healing; (e) neurological diseases e.g. cerebral anoxia and
 CC epilepsy; and (f) infectious diseases such as viral, bacterial, fungal
 CC and parasitic infections.
 CC Note: The sequence data for this patent did not form part of the
 CC printed specification, but was obtained in electronic format directly
 CC from WIPO at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 32220 BP; 7879 A; 8016 C; 7968 G; 8357 T; 0 other;
 Query Match 29.5%; Score 292.8; DB 22; Length 32220;
 Best Local Similarity 97.7%; Pred. No. 2.8e-62;
 Matches 297; Conservative 0; Mismatches 7; Indels 0; Gaps 0;
 QY 688 CAGTGGTCCGCCAGGTGGTACCAAGATGAGGAGTGGCAGGGTCCCTGGAGACC 747
 DB 9199 CACTGCTTTGTCAAGGTGGTGACCAAGATGAGGAGTGGCAGGGTCCCTGGAGACC 9258
 QY 748 CCATCTTGCCTTCCGCTCTACAGAGACCCCGCGCTGTGATGAGAGGATTTGGGG 807
 DB 9259 CCATCTTGCCTTCCGCTCTACAGAGACCCCGCGCTGTGATGAGAGGATTTGGGG 9318
 QY 808 GGTTCAGATAGGAATGGGGAGTGCAGAGCGCAAGCAGCAGCATGTAGAAATGAAC 867
 DB 9319 GGTTCAGATAGGAATGGGGAGTGCAGAGCGCAAGCAGCAGCATGTAGAAATGAAC 9378
 QY 868 CGTCCAGAGAGCCAGCAGCAGGAGACTGTCAGGGCCATCAGCGTGCATGTTGCTATT 927
 DB 9379 CGTCCAGAGAGCCAGCAGCAGGAGACTGTCAGGGCCATCAGCGTGCATGTTGCTATT 9438
 QY 928 GGAGTTTCATCAAAATGAGTGTGTTTGTAGTCTTGTGTCACAAAAAATAAAAAA 987
 DB 9439 GGAGTTTCATCAAAATGAGTGTGTTTGTAGTCTTGTGTCACAAAAAATAAAAAA 9498
 QY 988 AAAA 991
 DB 9499 AAAA 9502

RESULT 21
 AAX12008/c
 ID AAX12008 standard; DNA; 150 BP.
 XX
 AC AAX12008;
 XX
 DT 30-MAR-1999 (first entry)
 XX
 DE Human biallelic polymorphic DNA fragment EST376246b.
 XX
 KW Polymorphism; biallelic; human; forensic; paternity testing; disease;
 KW detection; phenotypic typing; characteristic; infection; hereditary;
 KW autoimmune disease; cancer; inflammation; drug; therapy; medicament;
 KW treatment; marker; ss.
 XX
 OS Homo sapiens.
 XX
 PN WO9820165-A2.
 XX
 PD 14-MAY-1998.
 XX
 PF 05-NOV-1997; 97WO-US20313.
 XX
 PR 06-NOV-1996; 96US-0030455.
 XX
 PA (WHEED) WHITEHEAD INST BIOMEDICAL RES.
 XX
 PI Hudson T, Lander ES, Wang D;
 XX
 DR WPI; 1998-286974/25.
 XX
 PT New isolated nucleic acid segments from the human genome - used for
 PT determining polymorphic forms for use in e.g. forensics, paternity
 PT testing or phenotypic typing for disease
 XX
 PS Claim 1; Page 212; 310pp; English.
 CC
 CC AAX10269-X12937 are human DNA fragments which contain biallelic
 CC polymorphic markers which have been isolated using the primers
 CC represented in AAX09121-X10268. The base occupying the polymorphic site
 CC is indicated by the appropriate IUPAC-RUB ambiguity code. These fragments
 CC can be used in methods for determining polymorphic forms in an individual
 CC for use in e.g. forensics, paternity testing or for phenotypic typing for
 CC diseases such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan
 CC syndrome, muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease,
 CC familial hypercholesterolemia, polycystic kidney disease, hereditary
 CC spherocytosis, von Willebrand's disease, tuberculous sclerosis, hereditary
 CC haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos
 CC syndrome, osteogenesis imperfecta, acute intermittent porphyria,
 CC autoimmune diseases, inflammation, cancer, diseases of the nervous
 CC system, infection by pathogenic microorganisms, and characteristics such
 CC as longevity, appearance (e.g. baldness, obesity), strength, speed,
 CC endurance, fertility, and susceptibility or receptivity to particular
 CC drugs or therapeutic treatments. The isolated polymorphic nucleic acid
 CC segments can also be used to produce medicaments for the treatment or
 CC prophylaxis of such diseases.
 XX
 SQ Sequence 150 BP; 30 A; 47 C; 32 G; 40 T; 1 other;
 Query Match 15.1%; Score 149.6; DB 19; Length 150;
 Best Local Similarity 99.3%; Pred. No. 1.6e-27;
 Matches 149; Conservative 1; Mismatches 0; Indels 0; Gaps 0;
 QY 820 GGAATGGGAGGTTCAGAGGACGCAAGCAGCAGCCATGTAGAAATGAACCGTCCAGAGAC 879
 DB 150 GGAATGGGAGGTTCAGAGGACGCAAGCAGCAGCCATGTAGAAATGAACCGTCCAGAGAC 91
 QY 880 CAAGCAGCGCAGAGGACTTGCAGGCCCATCAGCGTGCATGTTGCTATTGGAGTTTCATGCA 939
 DB 90 CAAGCAGCGCAGAGGACTTGCAGGCCCATCAGCGTGCATGTTGCTATTGGAGTTTCATGCA 91
 QY 940 AAATCAGTGTGTTTGTAGCTGCTCTTGGCCAC 969

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Db      30 AAATGAGTGTGTTTGTAGCTGCTCTTGCCAC 1
RESULT 23
AAAX12009/c
ID      AAX12009 standard; DNA; 150 BP.
AC      AAX12009;
XX
DT      30-MAR-1999 (first entry)
DE      Human biallelic polymorphic DNA fragment EST376246a.
XX
KW      Polymorphism; biallelic; human; forensic; paternity testing; disease;
KW      detection; phenotypic typing; characteristic; infection; hereditary;
KW      autoimmune disease; cancer; inflammation; drug; therapy; medication;
KW      treatment; marker; ss.
XX
OS      Homo sapiens.
XX
FN      WO9820165-A2.
XX
PD      14-MAY-1998.
XX
PF      05-NOV-1997; 97WO-US20313.
XX
PR      06-NOV-1996; 96US-0030455.
XX
PA      (WHED) WHITEHEAD INST BIOMEDICAL RES.
XX
PI      Hudson T, Lander ES, Wang D;
XX
DR      WPI; 1998-286974/25.
XX
PT      New isolated nucleic acid segments from the human genome - used for
PT      determining polymorphic forms for use in e.g. forensics, paternity
PT      testing or phenotypic typing for disease
XX
PS      Claim 1; Page 212; 310pp; English.
XX
CC      AAX10269-X12937 are human DNA fragments which contain biallelic
CC      polymorphic markers which have been isolated using the primers
CC      represented in AAX09121-X10268. The base occupying the polymorphic site
CC      is indicated by the appropriate IUPAC-IVB ambiguity code. These fragments
CC      can be used in methods for determining polymorphic forms in an individual
CC      for use in e.g. forensics, paternity testing or for phenotypic typing for
CC      diseases such as agammaglobulinemia, diabetes insipidus, Lesch-Nyhan
CC      syndrome, muscular dystrophy, Wiskott-Aldrich syndrome, Fabry's disease,
CC      familial hypercholesterolemia, polycystic kidney disease, hereditary
CC      spherocytosis, von Willebrand's disease, tuberculous sclerosis, hereditary
CC      haemorrhagic telangiectasia, familial colonic polyposis, Ehlers-Danlos
CC      syndrome, osteogenesis imperfecta, acute intermittent porphyria,
CC      autoimmune diseases, inflammation, cancer, diseases of the nervous
CC      system, infection by pathogenic microorganisms, and characteristics such
CC      as longevity, appearance (e.g. baldness, obesity), strength, speed,
CC      endurance, fertility, and susceptibility or receptivity to particular
CC      drugs or therapeutic treatments. The isolated polymorphic nucleic acid
CC      segments can also be used to produce medicaments for the treatment or
CC      prophylaxis of such diseases.
XX
SQ      Sequence 150 BP; 30 A; 46 C; 33 G; 40 T; 1 other;

Query Match      15.1%; Score 149.6; DB 19; Length 150;
Best Local Similarity 99.3%; Pred. No. 1.6e-27;
Matches 149; Conservative 1; Mismatches 0; Indels 0; Gaps 0;

QY      820 GGAATGGGAGGTGAGAGGACGAAAGCAGCAGCCATGTAGATGAACCGTCCAGAGGC 879
Db      150 GGAATGGGAGGTGAGAGGACGAAAGCAGCAGCCATGTAGATGAACCGTCCAGAGGC 91
QY      880 CAAGCAGGACGAGACTGCGAGGCCATCAGCGTCACTGTTGATTTGAGTTTCATGCA 939
Db      90 CAAGCAGGACGAGACTGCGAGGCCATCAGCGTCACTGTTGATTTGAGTTTCATGCA 31

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QY      940 AAATGAGTGTGTTTGTAGCTGCTCTTGCCAC 969
Db      30 AAATGAGTGTGTTTGTAGCTGCTCTTGCCAC 1

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RESULT 23
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ID      AEX97141 standard; cDNA; 1272 BP.
AC      AEX97141;
XX
DT      20-MAY-2003 (first entry)
DE      Human NOV93a cDNA.
XX
KW      NOVX; cytostatic; cardiant; antiarteriosclerotic; antiasthmatic; cancer;
KW      hypotensive; cardiomyopathy; bronchial asthma; gene therapy; vaccine;
KW      human; gene; ss.
XX
OS      Homo sapiens.
XX
FN      WO200272757-A2.
XX
PD      19-SEP-2002.
XX
PF      08-MAR-2002; 2002WO-US06908.
XX
PR      08-MAR-2001; 2001US-274101P.
PR      08-MAR-2001; 2001US-274194P.
PR      08-MAR-2001; 2001US-274281P.
PR      08-MAR-2001; 2001US-274322P.
PR      09-MAR-2001; 2001US-274849P.
PR      12-MAR-2001; 2001US-275235P.
PR      13-MAR-2001; 2001US-275578P.
PR      13-MAR-2001; 2001US-275579P.
PR      13-MAR-2001; 2001US-275601P.
PR      14-MAR-2001; 2001US-276000P.
PR      16-MAR-2001; 2001US-276776P.
PR      19-MAR-2001; 2001US-276994P.
PR      20-MAR-2001; 2001US-277239P.
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PR      27-MAR-2001; 2001US-278999P.
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PR      28-MAR-2001; 2001US-279344P.
PR      30-MAR-2001; 2001US-279995P.
PR      30-MAR-2001; 2001US-280233P.
PR      02-APR-2001; 2001US-280802P.
PR      02-APR-2001; 2001US-280822P.
PR      04-APR-2001; 2001US-280900P.
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PR      30-APR-2001; 2001US-283675P.
PR      03-MAY-2001; 2001US-287424P.
PR      03-MAY-2001; 2001US-288342P.
PR      15-MAY-2001; 2001US-288528P.
PR      16-MAY-2001; 2001US-291130P.
PR      16-MAY-2001; 2001US-291099P.
PR      30-MAY-2001; 2001US-291240P.
PR      31-MAY-2001; 2001US-294885P.
PR      31-MAY-2001; 2001US-294889P.
PR      18-JUN-2001; 2001US-299027P.
PR      19-JUN-2001; 2001US-299303P.
PR      19-JUN-2001; 2001US-299310P.
PR      10-JUL-2001; 2001US-304354P.
PR      31-JUL-2001; 2001US-309198P.

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PR 16-AUG-2001; 2001US-312903P.
 PR 10-SEP-2001; 2001US-318462P.
 PR 12-SEP-2001; 2001US-318770P.
 PR 27-SEP-2001; 2001US-325430P.
 PR 27-SEP-2001; 2001US-325681P.
 PR 18-OCT-2001; 2001US-330380P.
 PR 31-OCT-2001; 2001US-335301P.
 PR 14-NOV-2001; 2001US-332172P.
 PR 14-NOV-2001; 2001US-332271P.
 PR 14-NOV-2001; 2001US-332272P.
 PR 14-NOV-2001; 2001US-333184P.
 PR 14-NOV-2001; 2001US-333272P.
 PR 21-NOV-2001; 2001US-332094P.
 PR 03-DEC-2001; 2001US-337426P.
 PR 03-DEC-2001; 2001US-338092P.
 PR 04-DEC-2001; 2001US-337185P.
 PR 03-JAN-2002; 2002US-345705P.
 PR 07-MAR-2002; 2002US-0092900.
 XX
 PA (CURA-) CURAGEN CORP.
 XX
 XX Padigar M, Spytek KA, Shenoy SG, Taupier RJ, Pena CEA, Li L;
 PI Zerhusen BD, Gusev V, Ji W, Gorman L, Miller CE, Kekuda R;
 PI Patturajan M, Gangolli E, Vernet CAM, Guo X, Tchernev V;
 PI Fernandes ER, Casman SJ, Malyankar UM, Gerlach V, Liu Y;
 PI Anderson D, Spaderna SK, Catterton E, Burgess C, Leite M, Zhong H;
 PI Alsobrook JP, Lepley DM, Rieger DK;
 XX
 XX WPI; 2002-723332/78.
 DR P-PSDB; ABU65174.
 XX
 XX NOVX polypeptides and polynucleotides, useful for preventing or
 PT treating a disorder associated with aberrant NOVX expression or
 PT activity e.g., cancer, hypertension, atherosclerosis, cardiomyopathy or
 PT bronchial asthma
 XX
 XX Claim 13; Page 368-369; 1103pp; English.
 PS
 XX This invention describes novel human NOVX polypeptides which have
 CC cytostatic, cardiant, antiarteriosclerotic, antiasthmatic and
 CC hypotensive activity. Pharmaceutical compositions comprising the NOVX
 CC proteins or nucleic acid molecules or NOVX antibodies are useful for
 CC preventing or treating a disorder associated with aberrant NOVX
 CC expression or activity e.g. cancer, hypertension, atherosclerosis,
 CC cardiomyopathy or bronchial asthma. The products of the invention can
 CC be used for gene therapy or in a vaccine ABX97008-ABX97185 are cDNA
 CC fragments amplified and isolated by the PCR primers and probes
 CC represented in ABX13460-ABX13462 and ABX97186-ABX97593. ABX97008-ABX97185
 CC encode the NOVX proteins described in ABU65041-ABU65218.
 XX
 XX Sequence 1272 BP; 182 A; 521 C; 356 G; 213 T; 0 other;
 SQ
 Query Match 4.8%; Score 47.2; DB 24; Length 1272;
 Best Local Similarity 48.8%; Pred. No. 0.084;
 Matches 127; Conservative 0; Mismatches 133; Indels 0; Gaps 0;
 QY 256 CCAAGAACATCAAGTGGCCAGAGAGGTGGTGAAGACCCAGCGCGCTCTTCAACC 315
 Db 711 CCCCAGACAGAGCTGATCGTGGTGGCGGAACCCCGTGACCCCGGCCACTCCGACT 770
 QY 316 TCAAGCTCACCTCAAGTCCAGTCCAGACCTGCTCAGCTTCTGCGGGGCTCTCCA 375
 Db 771 AGGCCCCAGACGCTCTCCAGACCCCGGGCTCCAGCTTCCGCGCTTGGCTTCGGCC 830
 QY 376 CTTAGGTGCTGAGTGGAGACGTCAGCGCTACAGATGACATGGAGCTGTGGTCAAGC 435
 Db 831 ACGGCTGCGGCCCGCTGGACACAGCTGGAGCGCGTCCGATCGCTGTACGCCGACG 890
 QY 436 CAGTGTCTGAGCTGGGGCCACTTCACTTCAGACAGAGGGCGAGGCCCGGAGGTGG 495
 Db 891 ACCTTGACCACTGGCTGGCGTACTTCCCGCTTCCCACTTCTGTTGTCAGCGGGAGC 950
 QY 496 AGATGATCTGCCAGGCGTCC 515

Db 951 GTCTGGTCAGCGACCCGCGCC 970
 RESULT 24
 ABQ76896
 ID ABQ76896 standard; DNA; 53226 BP.
 XX
 AC ABQ76896;
 XX
 DT 13-MAR-2003 (first entry)
 XX
 DE Human G-protein coupled receptor DNA SEQ ID 3.
 XX
 KW G-protein coupled receptor; secretin receptor subfamily; human; SNP;
 KW GPCR; protease; Parkinson's disease; gene; chromosome X;
 KW single nucleotide polymorphism; ds.
 XX
 OS Homo sapiens.
 XX
 XX Key Location/Qualifiers
 FT CDS 3000..50651
 FT /*tag= a
 FT /product= "GPCr"
 FT /note "this coding sequence is interrupted by
 FT 13 introns"
 FT variation replace (1745,c)
 FT /*tag= b
 FT /note= "SNP, single nucleotide polymorphism"
 FT replace (1755,g)
 FT /*tag= c
 FT /note= "SNP, single nucleotide polymorphism"
 FT replace (1961,g)
 FT /*tag= d
 FT /note= "SNP, single nucleotide polymorphism"
 FT 3000..3088
 FT /*tag= e
 FT /number= 1
 FT 3089..3874
 FT /*tag= f
 FT /number= 1
 FT 3875..4038
 FT /*tag= g
 FT /number= 2
 FT 4039..6037
 FT /*tag= h
 FT /number= 2
 FT 5411.g
 FT /*tag= i
 FT /note= "SNP, single nucleotide polymorphism"
 FT replace (5760,a)
 FT /*tag= j
 FT /note= "SNP, single nucleotide polymorphism"
 FT 6038..6170
 FT /*tag= k
 FT /number= 3
 FT 6171..8059
 FT /*tag= l
 FT /number= 3
 FT 8060..8178
 FT /*tag= m
 FT /number= 4
 FT 8179..15910
 FT /*tag= n
 FT /number= 4
 FT replace (11390,c)
 FT /*tag= o
 FT /note= "SNP, single nucleotide polymorphism"
 FT replace (11390,t)
 FT /*tag= p
 FT /note= "SNP, single nucleotide polymorphism"
 FT 15911..16127
 FT /*tag= q


```
FT intron /number= 5
FT 16128..17484
FT /tag= r
FT /number= 5
FT variation replace (16988,a)
FT /tag= s
FT /note= "SNP, single nucleotide polymorphism"
FT exon 17485..17647
FT /tag= t
FT /number= 6
FT intron 17648..32332
FT /tag= u
FT /number= 6
FT variation replace (18361,g)
FT /tag= v
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (19769,c)
FT /tag= w
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (22910,t)
FT /tag= x
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (22935,t)
FT /tag= y
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (24206,g)
FT /tag= z
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (24774,a)
FT /tag= aa
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (24869,c)
FT /tag= ab
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (25768,a)
FT /tag= ac
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (26697,t)
FT /tag= ad
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (26697,c)
FT /tag= ae
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (28359,c)
FT /tag= af
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (28470..28471,t)
FT /tag= ag
FT /note= "a single nucleotide polymorphism (SNP) can
FT result in a deletion at this position"
FT variation replace (29781,g)
FT /tag= ah
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (30182,a)
FT /tag= ai
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (31772,t)
FT /tag= aj
FT /note= "SNP, single nucleotide polymorphism"
FT variation replace (31936,g)
FT /tag= ak
FT /note= "SNP, single nucleotide polymorphism"
FT exon 32333..32467
FT /tag= al
FT /number= 7
FT intron 32468..36825
FT /tag= am
FT /number= 7
FT exon 36826..36948
FT /tag= an
FT /product= 8
FT 36949..38129
FT /tag= ao
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```
FT exon /number= 8
FT 38130..38175
FT /tag= ap
FT /number= 9
FT intron 38176..39984
FT /tag= aq
FT /number= 9
FT exon 39985..40088
FT /tag= ar
FT /number= 10
FT intron 40089..42455
FT /tag= as
FT /number= 10
FT exon 42456..42577
FT /tag= at
FT /number= 11
FT intron 42578..44422
FT /tag= au
FT /number= 11
FT variation replace (42767..42767,c)
FT /tag= av
FT /note= "a single nucleotide polymorphism (SNP) can
FT result in a deletion at this position"
FT exon 44423..44691
FT /tag= aw
FT /number= 12
FT intron 44692..47818
FT /tag= ax
FT /number= 12
FT exon 47819..47897
FT /tag= ay
FT /number= 13
FT intron 47898..50266
FT /tag= az
FT /number= 13
FT variation replace (48839,t)
FT /tag= ba
FT /note= "SNP, single nucleotide polymorphism"
FT exon 50267..50651
FT /tag= bb
FT /number= 14
FT variation replace (52265,g)
FT /tag= bc
FT /note= "SNP, single nucleotide polymorphism"
XX US2002142951-A1.
XX 03-OCT-2002.
XX 28-MAR-2001; 2001US-0818264.
XX 28-MAR-2001; 2001US-0818264.
XX (WEBS/) WEBSTER M.
XX (BEAS/) BEASLEY E M.
XX (KETCH/) KETCHUM K A.
XX (DFRA/) DI FRANCESCO V.
XX Webster M, Beasley EM, Ketchum KA, Di Francesco V;
```

Query Match 4.7%; Score 47; DB 25; Length 53226;
Best local similarity 78.9%; Pred. No. 0.33;
Matches 56; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 922 GTATTTGGAGTTCATGCAAAATGAGTGTGTTTGTAGTCTTGGCCACAAAAA 981

Db 29825 GTTTTAGATTTCATGCTAAATGATAGTTTAGTCTTGGCCACAAAAA 29884

QY 982 AAAAAAAAAA 992

Db 29885 AAAAGAGGTAA 29895

RESULT 25
 AAX83003/C
 ID AAX83003 standard; DNA; 87350 BP.
 AC AAX83003;
 XX
 XX 31-AUG-1999 (first entry)
 XX
 DE Human WRN genomic sequence.
 XX
 KW Human; WRN; Werner's syndrome; detection; diagnosis; autosomal;
 KW recessive disorder; phenotype; ss.
 XX
 OS Homo sapiens.
 XX
 XX WO9724435-A1.
 PN 10-JUL-1997.
 XX
 XX 30-DEC-1996; 96WO-US20785.
 PF
 XX 12-APR-1996; 96US-0632175.
 PR 29-DEC-1995; 95US-0009409.
 PR 29-DEC-1995; 95US-0580539.
 PR 30-JAN-1996; 96US-0010835.
 PR 30-JAN-1996; 96US-0594242.
 XX
 PA (DARW-) DARWIN MOLECULAR CORP.
 PA (OSHI/) OSHIMA J.
 XX
 PI Fu Y, Mulligan J, Oshima J, Schellenberg GD, Yu C;
 XX
 DR WPI; 1997-363671/33.
 XX
 XX Isolated nucleic acid molecule encoding the WRN gene product -
 PT useful for detection and treatment of Werner's syndrome, and related
 PT diseases
 PT
 XX
 PS Claim 1; Fig 5A-U; 153pp; English.
 XX
 CC This sequence represents the genomic region containing the coding
 CC sequence for the human WRN gene which encodes a protein related to
 CC Werner's syndrome. The products can be used for the detection and
 CC treatment of Werner's syndrome (WS), an autosomal recessive disorder
 CC with a complex phenotype, as well as related diseases.
 CC
 XX Sequence 87350 BP; 25621 A; 16221 C; 17012 G; 28450 T; 46 other;
 SQ
 Query Match 4.7%; Score 47; DB 18; Length 87350;
 Best Local Similarity 78.9%; Pred. No. 0.39;
 Matches 56; Conservative 0; Mismatches 15; Indels 0; Gaps 0;
 QY 922 GTATTGGAGTTCATGCAAAATGAGTGCTGTTTGTAGCTGCTTGGCCACAAAAA 981
 DB 18894 GTATTGGAGTTCCTGCTAAATGAGTGGATTTCAGCTGCTTTGGCACAACAAAAA 18835
 QY 982 AAAAAA 992
 DB 18834 ATGGTTAACAA 18824
 RESULT 26
 AAS68336/C
 ID AAS68336 standard; cDNA; 636 BP.
 AC AAS68336;
 XX
 XX 13-FEB-2002 (first entry)
 XX
 DE DNA encoding novel human diagnostic protein #4140.
 XX
 KW Human; chromosome mapping; gene mapping; gene therapy; forensic;

KW food supplement; medical imaging; diagnostic; genetic disorder; ss.
 XX
 OS Homo sapiens.
 PN WO200175067-A2.
 XX
 PD 11-OCT-2001.
 XX
 XX 30-MAR-2001; 2001WO-US08631.
 XX
 PF 31-MAR-2000; 2000US-0540217.
 PR 23-AUG-2000; 2000US-0649167.
 XX
 XX (HYSE-) HYSEQ INC.
 PA
 XX Drmanac RT, Liu C, Tang YT;
 PI WPI; 2001-639362/73.
 XX P-PSDB; ABG04149.
 DR
 DR
 XX
 XX New isolated polynucleotide and encoded polypeptides, useful in
 PT diagnostics, forensics, gene mapping, identification of mutations
 PT responsible for genetic disorders or other traits and to assess
 PT biodiversity -
 XX
 XX Claim 1; SEQ ID No 4140; 103pp; English.
 XX
 CC The invention relates to isolated polynucleotide (I) and
 CC polypeptide (II) sequences. (I) is useful as hybridisation probes,
 CC polymerase chain reaction (PCR) primers, oligomers, and for chromosome
 CC and gene mapping, and in recombinant production of (II). The
 CC polynucleotides are also used in diagnostics as expressed sequence tags
 CC for identifying expressed genes. (I) is useful in gene therapy techniques
 CC to restore normal activity of (II) or to treat disease states involving
 CC (II). (II) is useful for generating antibodies against it, detecting or
 CC quantitating a polypeptide in tissue, as molecular weight markers and as
 CC imaging of sites expressing (II). (I) and (II) are useful for treating
 CC disorders involving aberrant protein expression or biological activity.
 CC The polypeptide and polynucleotide sequences have applications in
 CC diagnostics, forensics, gene mapping, identification of mutations
 CC responsible for genetic disorders or other traits to assess biodiversity
 CC and to produce other types of data and products dependent on DNA and
 CC amino acid sequences. AAS64197-AAS94564 represent novel human
 CC diagnostic coding sequences of the invention.
 CC Note: The sequence data for this patent did not appear in the printed
 CC specification, but was obtained in electronic format directly from WIPO
 CC at ftp.wipo.int/pub/published_pct_sequences.
 XX
 SQ Sequence 636 BP; 103 A; 246 C; 180 G; 107 T; 0 other;
 Query Match 4.5%; Score 44.4; DB 23; Length 636;
 Best Local Similarity 88.9%; Pred. No. 0.33;
 Matches 48; Conservative 0; Mismatches 6; Indels 0; Gaps 0;
 QY 57 CGAGCATGGGGCTCCCTGGGCTGTTCTGCTTGGCCGCTGCTGCACGACC 110
 DB 263 CGAGCATGGGGCTCCCTGGGCTGTTCTGCTTGGCCGCTGCTGCACGACC 210
 RESULT 27
 ABZ80229
 ID ABZ80229 standard; DNA; 249999 BP.
 XX
 XX AC ABZ80229;
 XX
 XX 02-JUN-2003 (first entry)
 XX
 DE Human transdormin gene region genomic DNA SEQ ID NO:26.
 KW Neuroprotective; nootropic; cerebroprotective; analgesic; gene therapy;
 KW central nervous system disorder; CNS disorder; multiple sclerosis;
 KW nerve injury; neuropathic pain; stroke; trauma; non-CNS disorder; transdormin;

KW tramodrin; human; chromosome 5; gene; ds.
OS Homo sapiens.
XX WO2003016502-A2.
PN 27-FEB-2003.
XX 21-AUG-2002; 2002WO-US26637.
PF 21-AUG-2001; 2001US-313907P.
XX 21-AUG-2002; 2002US-0225810.
PR (MCLA-) MCLAUGHLIN RES INST.
XX Birmingham JR;
PI WPI; 2003-278567/27.
DR New nucleic acid sequence encoding tramodrin, e.g. mouse tramd 1, mouse tramd 2, mouse tramd 3, human tramd 1, human tramd 2, human tramd 3 or rat tramd 1, useful for treating CNS, e.g. stroke, multiple sclerosis, trauma, neuropathic pain -
XX Example 6; Fig 9; 177pp; English.
PS The present invention describes an isolated nucleic acid sequence comprising a cDNA sequence encoding mouse tramodrin (tramd) 2, mouse tramd 3, human tramd 1, human tramd 2, human tramd 3 or rat tramd 1, or the genomic sequence of mouse tramd 1 or mouse tramd 3. Mouse tramd 1 is located to chromosome 11, whereas human tramd 1 is located to chromosome 5q31-33. The tramd sequences have neuroprotective, nootropic, analgesic and cerebroprotective activities, and can be used in gene therapy. The nucleic acid sequences are useful for diagnosing and treating central nervous system (CNS) disorders such as multiple sclerosis, nerve injury, neuropathic pain, stroke or trauma, and non-CNS disorders. The present sequence represents the genomic sequence of the human tramodrin gene region, which is given in the exemplification of the present invention.
XX Sequence 249999 BP; 75050 A; 54012 C; 51931 G; 67805 T; 1201 other;
SQ Query Match 4.4%; Score 44; DB 25; Length 249999;
Best Local Similarity 77.9%; Pred. No. 3.1;
Matches 53; Conservative 0; Mismatches 15; Indels 0; Gaps 0;
QY 922 GTATTGGAGTTCATGCAAAATGCTGTCTTTAGCTGCTCTTGCACAAAAA 981
Db 75447 GTATTAAAATCTGTGAATGAGTAAATTTAGCTGCTCTTGCACAAAAA 75506
QY 982 AAAAAAAA 989
Db 75507 CCAATAA 75514
RESULT 28
ABT07673
ID ABT07673 standard; cDNA; 560 BP.
AC ABT07673;
XX 14-NOV-2002 (first entry)
XX Human breast cancer associated coding sequence SEQ ID NO: 152.
DE Human; breast specific gene; breast specific protein; breast cancer;
KW gene therapy; cytostatic; gene; ss.
XX Homo sapiens.
OS WO200264611-A1.
XX 22-AUG-2002.
PD

PF 12-FEB-2002; 2002WO-US04197.
XX 13-FEB-2001; 2001US-268292P.
XX (DIAD-) DIADEXUS INC.
PI Salceda S, Macina RA, Hu P, Recipon H, Karra K, Cafferkey R;
PI Sun Y, Liu C;
XX WPI; 2002-657582/70.
XX New breast specific nucleic acids and proteins, useful for identifying, diagnosing, monitoring, staging, imaging, and treating breast cancer and non-cancerous disease states in breast tissue, and in gene therapy -
XX Claim 1; Page 269; 367pp; English.
XX The present invention provides human breast specific coding sequences and proteins. These can be used in the diagnosis and treatment of breast cancer and non-cancerous diseases of the breast. The present sequence is a coding sequence of the invention.
XX Sequence 560 BP; 173 A; 109 C; 139 G; 139 T; 0 other;
SQ Query Match 4.4%; Score 43.4; DB 24; Length 560;
Best Local Similarity 76.8%; Pred. No. 0.56;
Matches 53; Conservative 0; Mismatches 16; Indels 0; Gaps 0;
QY 924 ATTTGGAGTTCATGCAAAATGAGTGTGTTTGTAGCTGCTCTTGCACAAAAA 983
Db 408 ATATGGGATTCATCTACTAATGAGTAGATTGCTTCTTGGCACAATAACAAAA 467
QY 984 AAAAAAAA 992
Db 468 ATGCCACAA 476
RESULT 29
ABT07672
ID ABT07672 standard; cDNA; 509 BP.
AC ABT07672;
XX 14-NOV-2002 (first entry)
XX Human breast cancer associated coding sequence SEQ ID NO: 151.
DE Human; breast specific gene; breast specific protein; breast cancer;
KW gene therapy; cytostatic; gene; ss.
XX Homo sapiens.
OS WO200264611-A1.
XX 22-AUG-2002.
PD 12-FEB-2002; 2002WO-US04197.
XX 13-FEB-2001; 2001US-268292P.
PR (DIAD-) DIADEXUS INC.
PI Salceda S, Macina RA, Hu P, Recipon H, Karra K, Cafferkey R;
PI Sun Y, Liu C;
XX WPI; 2002-657582/70.
XX New breast specific nucleic acids and proteins, useful for identifying, diagnosing, monitoring, staging, imaging, and treating breast cancer and non-cancerous disease states in breast tissue, and in gene therapy -
XX

PS Claim 1; Page 268-269; 367pp; English.

CC The present invention provides human breast specific coding sequences and

CC proteins. These can be used in the diagnosis and treatment of breast

CC cancer and non-cancerous diseases of the breast. The present sequence is

CC a coding sequence of the invention.

XX

SQ Sequence 509 BP; 151 A; 109 C; 123 G; 126 T; 0 other;

Query Match 4.3%; Score 43; DB 24; Length 509;

Best Local Similarity 77.6%; Pred. No. 0.66;

Matches 52; Conservative 0; Mismatches 15; Indels 0; Gaps 0;

QY 926 TTGGAGTTCATGCACAAATGAGTGTGTTTAGTCTCTTCCACAAAAAATAAAAAA 985

Db 359 TTGGGATTCATCTAATGAGTAGATTAGTCTCTTCCACAAATAACAAAAAAT 418

QY 986 AAAAAA 992

Db 419 GCCACAA 425

RESULT 30

AA138336/c

ID AAL38336 standard; DNA; 143899 BP.

XX

AC AAL38336;

XX

DT 15-AUG-2002 (first entry)

XX

DE Genomic sequence encoding a human Ngr2 protein.

XX

KW Cerebroprotective; neuroprotective; cytostatic; Nogo receptor homologue;

KW Ngr2; Ngr3; axonal growth; central nervous system; CNS; cerebral injury;

KW spinal cord injury; stroke; demyelinating disease; multiple sclerosis;

KW monophasic demyelination; encephalomyelitis; Marchiafava-Bignami disease;

KW multifocal leukoencephalopathy; panencephalitis; Spongy degeneration;

KW Alexander's disease; Canavan's disease; metachromatic leukodystrophy;

KW Krabbe's disease; immune; bait protein; genetic mapping; gene therapy;

KW transgenic animal; unregulated cellular growth; cancer; tumour; human;

KW gene; ds.

XX

OS Homo sapiens.

XX

PN WO200229059-A2.

XX

PD 11-APR-2002.

XX

PF 06-OCT-2001; 2001WO-US31488.

XX

PR 06-OCT-2000; 2000US-238361P.

XX

PA (UYVA) UNIV YALE.

PA (BIOU) BIOGEN INC.

XX

PI Strittmatter SM, Cate RL, Sah DMW;

XX

DR WPI; 2002-416677/44.

XX

PT Novel Nogo receptor homolog polypeptide, Ngr2 or Ngr3, useful for

PT treating central nervous system disorder, cerebral injury, spinal cord

PT injury, stroke, and demyelinating diseases -

XX

PS Example 2; Page 176-214; 277pp; English.

XX

CC The invention relates to a Nogo receptor homologue polypeptide, Ngr2 or

CC Ngr3, comprising a 50 amino acid LRRCT sequence, a 284 amino acid NTLRRCT

CC sequence, or a 420, 461 or 392 amino acid sequence, all given in the

CC specification. The Ngr3 protein or its binding antibody is useful for

CC decreasing inhibition of axonal growth of a central nervous system (CNS)

CC neuron, by contacting the neuron Ngr3 or its antibody, and for treating

CC CNS disease, disorder or injury. Ngr3 or a vector comprising Ngr3 is

CC useful for treating cerebral injury, spinal cord injury, stroke,

CC demyelinating diseases, e.g. multiple sclerosis, monophasic

CC demyelination, encephalomyelitis, multifocal leukoencephalopathy,

CC panencephalitis, Marchiafava-Bignami disease, Spongy degeneration,

CC Alexander's disease, Canavan's disease, metachromatic leukodystrophy and

CC Krabbe's disease. Ngr3 is useful for inducing an immune response in a

CC mammal against Ngr3, as a bait protein in a two-hybrid or three-hybrid

CC assay, and as a research tool for identification, characterisation and

CC purification of interacting, regulatory proteins. The nucleotide

CC sequences of the invention are useful for screening for RFP associated

CC with certain disorders, for genetic mapping, and for gene therapy. The

CC vector containing Ngr3 is useful for producing non-human transgenic

CC animals. The Ngr3 binding antibody is useful for isolating and purifying

CC Ngr3, for localisation and/or quantitation of Ngr3, and for diagnostic

CC and therapeutic purposes. The sequences of the invention, vectors and

CC antibodies are useful for treating or preventing unregulated cellular

CC growth such as cancer and tumour growth. This polynucleotide sequence

CC represents the genomic sequence encoding a human Ngr2 protein of the

CC invention..

XX

SQ Sequence 143899 BP; 36346 A; 35277 C; 35318 G; 35657 T; 1301 other;

Query Match 4.3%; Score 42.8; DB 24; Length 143899;

Best Local Similarity 56.1%; Pred. No. 5;

Matches 101; Conservative 0; Mismatches 77; Indels 2; Gaps 1;

QY 804 GGGGGTTTCAGCATAGGGAATGGGAGGTTCAGAGACGCAAGCAGCAGCCATGTAGAT 863

Db 92710 GGGTGGAGAGGAGATGGGAGATGCAGGTGCAAGGACAAAAAGGAGCAGATGAGCAGGAT 92651

QY 864 GAACCGTTCAGAGAGCCAGGACGCGAGGACTGTCAGCGCCATCAGCGTGCATCTTCGT 923

Db 92650 GAACAGGCTAGAGATCTAATCCAGCAGCAGAGACTGTAGTCTATCAACTGCATTGTG 92591

QY 924 ATTTGGAGTTCATGCAAAATAGTGTGTTTATGCTGCTTTCGCCCAAAAAAATAAAAA 983

Db 92590 TTTGGGACTT--TCCTAAACAAGTAGATTTTAACTGCTTCTGTACAAAAACAGGTAGAAA 92533

RESULT 31

AAF21712/c

ID AAF21712 standard; DNA; 752 BP.

XX

AC AAF21712;

XX

DT 27-MAR-2001 (first entry)

XX

DE Human breast and ovarian cancer associated antigen gene SEQ ID 99.

XX

KW Human; breast cancer; ovarian cancer; cytostatic; immunosuppressive;

KW neurotropic; neuroprotective; antiviral; antiallergic; hepatotropic;

KW antidiabetic; antiinflammatory; antiulcer; vulnery; anticonvulsant;

KW antibacterial; antifungal; antiparasitic; cardiant; immune disorder;

KW Addison's disease; allergy; autoimmune haemolytic anaemia;

KW autoimmune thyroiditis; diabetes mellitus; Crohn's disease;

KW multiple sclerosis; rheumatoid arthritis; ulcerative colitis;

KW cardiovascular disorder; wound healing; neurological disease; ds.

XX

OS Homo sapiens.

XX

PN WO200055173-A1.

XX

PD 21-SEP-2000.

XX

PF 08-MAR-2000; 2000WO-US05881.

XX

PR 12-MAR-1999; 99US-0124270.

XX

PA (HUMA-) HUMAN GENOME SCI INC.

XX

PI Rosen CA, Ruben SM;

XX

DR WPI; 2000-611515/58.

DR P-PSDB; AAB59809.

XX New human breast and ovarian cancer associated gene sequences and the
PT polypeptides encoded by these genes, useful in the prevention,
PT treatment and diagnosis of cancer, immune disorders, cardiovascular
PT disorders and neurological diseases -
XX
PS Claim 1; Page 555-556; 1299pp; English.
XX
XX Sequences AAF21614 - AAF22031 represent DNA sequences encoding human
CC proteins AAB59711 - AAB59128. The DNA and protein sequences are
CC associated with breast and ovarian cancer. Included in the invention are
CC sequences AAF22032 - AAF22040 and AAB59129 which are used in the
CC isolation and characterisation of the DNA and protein sequences of the
CC invention. The breast and ovarian cancer associated DNA, protein, agonist
CC or antagonist sequences exhibit cytostatic; immunosuppressive;
CC neurotropic; neuroprotective; antiviral; antiallergic; hepatotropic;
CC antidiabetic; antiinflammatory; antiulcer; vulnerary; anticonvulsant;
CC antibacterial; antifungal; antiparasitic and cardiant activity. The
CC polynucleotide and protein sequences are used in the diagnosis of cancer,
CC particularly breast and ovarian cancer. The nucleic acid sequences,
CC proteins, agonists and agonists may also be used in the diagnosis,
CC prevention and treatment of immune disorders e.g. Addison's disease,
CC allergies, autoimmune haemolytic anaemia, autoimmune thyroiditis,
CC diabetes mellitus, Crohn's disease, multiple sclerosis, rheumatoid
CC arthritis and ulcerative colitis; cardiovascular disorders such as
CC myocardial ischaemias; wound healing; neurological diseases such as
CC cerebral anoxia and epilepsy; and infectious diseases.
XX
SQ Sequence 752 BP; 149 A; 275 C; 219 G; 104 T; 5 other;
Query Match 4.2%; Score 41.8; DB 21; Length 752;
Best Local Similarity 51.3%; Pred. NO. 1.5;
Matches 97; Conservative 0; Mismatches 92; Indels 0; Gaps 0;
QY 410 GATGCACTGGAGCTGTGTTCCAGCCAGTGTCTGAGCTGGGGCCCACTTCACTCTGCA 469
Db 260 GCTGGCTGGCAGATGCAGTCCAGCGGCTGTGTAGCGCAGGACGCTGGCCAGACC 201
QY 470 GGA CAGAGGGGCGAGCCGCCAGGGTGGAGATGATCTGCCAGCGCTCTGGGCGAGCCACC 529
Db 200 CCGCAGAGGGGGCGCCCGCAGACCGCGCGCCCTGCGCGTGTGCGCAGTCAGCACC 141
QY 530 TATCACAACAGCCTGTGTCGGGAGGATGGCAGGTCCACTGCAGCAGACCATGCCA 589
Db 140 TGCACAGAACCCGCTTATGGCCATCGCGCTTGGGGGGCGAGTAGCGGCTCTGCACCGA 81
QY 590 CAGGCAGCC 598
Db 80 CAGGGAGGC 72
RESULT 32
AAC77293/c
ID AAC77293 standard; cDNA; 1417 BP.
XX
AC AAC77293;
XX
DT 08-FEB-2001 (first entry)
XX
DE Human ORFX ORF2848 polynucleotide sequence SEQ ID NO:5695.
XX
XX Human; open reading frame; ORFX; detection; cytostatic; hepatotropic;
KW vulnerary; antiparasitic; antiparkinsonian; neurotropic; neuroprotective;
KW anticonvulsant; osteopathic; antiarthritic; immunosuppressant; cardiant;
KW immunostimulant; thrombolytic; coagulant; vasotropic; antidiabetic;
KW hypotensive; dermatological; immunosuppressive; antiinflammatory;
KW antiviral; antibacterial; antifungal; antirheumatic; antithyroid;
KW antianemic; gene therapy; cancer; proliferative disorder; hypertension;
KW neurodegenerative disease; osteoarthritis; graft vs host disease;
KW cardiovascular disease; diabetes mellitus; hypothyroidism; SCID; AIDS;
KW cholesterol ester storage; systemic lupus erythematosus; infection;
KW severe combined immunodeficiency; malaria; autoimmune disorder; asthma;
KW allergy; aplastic anaemia; nocturnal haemoglobinuria; burn; wound;

KW bone damage; cartilage damage; antiinflammatory disease; coagulation;
KW thrombosis; contraceptive; ss.
XX Homo sapiens.
XX WO200058473-A2.
XX 05-OCT-2000.
PD
XX 31-MAR-2000; 2000WO-US08621.
PF
XX 31-MAR-1999; 99US-0127607.
PR 02-APR-1999; 99US-0127636.
PR 05-APR-1999; 99US-0127728.
PR 30-MAR-2000; 2000US-0540763.
XX (CURA-) CURAGEN CORP.
PA Shimkets RA, Leach M;
PI WPI; 2000-602362/57.
XX P-PSDB; AAB43084.
DR
DR
DR
XX Novel nucleic acids and peptides derived from open reading frame X,
PT useful for treating e.g. cancers, proliferative disorders,
PT neurodegenerative disorders and cardiovascular disease -
XX
PS Claim 5; Page 4858-4859; 5507pp; English.
XX
CC AAC74446 to AAC77606 encode the proteins given in AAB40237 to AAB43397,
CC which represent the human ORFX open reading frames 1 to 3161. The ORFX
CC sequences have activities such as: cytostatic; hepatotropic; vulnerary;
CC antiparasitic; antiparkinsonian; neurotropic; neuroprotective;
CC osteopathic; anticonvulsant; antiarthritic; immunosuppressant;
CC immunostimulant; cardiant; thrombolytic; coagulant; vasotropic;
CC antidiabetic; hypotensive; dermatological; immunosuppressive;
CC antiinflammatory; antibacterial; antiviral; antifungal; antirheumatic;
CC antithyroid; and antianemic. The sequences can be used for determining
CC the presence of or predisposition to, or preventing or treating
CC pathological conditions associated with an ORFX-associated disorder. The
CC nucleic acids can be used to express ORFX proteins in gene therapy
CC vectors. The proteins and nucleic acids may be used to treat cancers,
CC proliferative disorders, neurodegenerative disorders, osteoarthritis,
CC graft vs host disease, cardiovascular disease, diabetes mellitus,
CC hypertension, hypothyroidism, cholesterol ester storage, systemic lupus
CC erythematosus, severe combined immunodeficiency (SCID), AIDS, viral,
CC bacterial or fungal infection, malaria, autoimmune disorders, asthma,
CC allergies, aplastic anaemia, burns, wounds, bone and cartilage damage,
CC nocturnal haemoglobinuria, antiinflammatory disease; to enhance
CC coagulation; to inhibit thrombosis; and as a contraceptive.
XX
SQ Sequence 1417 BP; 211 A; 499 C; 473 G; 234 T; 0 other;
Query Match 4.2%; Score 41.8; DB 21; Length 1417;
Best Local Similarity 51.3%; Pred. NO. 1.9;
Matches 97; Conservative 0; Mismatches 92; Indels 0; Gaps 0;
QY 410 GATGCACTGGAGCTGTGTTCCAGCCAGTGTCTGAGCTGGGGCCCACTTCACTCTGCA 469
Db 987 GCTGGGCTGGCAGATGCAGTCCAGCGGCTGTGTAGCGCAGGACGCTGGCCAGACC 928
QY 470 GGA CAGAGGGGCGAGCCGCCAGGGTGGAGATGATCTGCCAGGGTCTCTGGGGCGAGCCACC 529
Db 927 CCGCAGAGGGGGCGCCCGCAGACCGCGCGCCCTGCCCGTAGTCGCCAGTCAGCACC 868
QY 530 TATCACAACAGCCTGTATCGGAGGATGGGAGGTCCACCTGCAGCAGACCATGCCA 599
Db 867 TGCACAGAACCCGCTTATGGCCATCGCGCTTGGGGGGCGAGTAGCGGCTCTGCACCGA 808
QY 590 CAGGCAGCC 598
Db 807 CAGGGAGGC 799

RESULT 33
AAC77637
ID AAC77637 standard; cDNA; 2059 BP.
XX
AC AAC77637;
XX
DT 08-FEB-2001 (first entry)
XX
DE Human cancer associated gene sequence SEQ ID NO:31.
XX
KW Human; cancer associated gene; cancer antigen; detection; cancer;
KW diagnosis; cytostatic; proliferative; vulnery; immunomodulator;
KW antidiabetic; antiasthmatic; antirheumatic; antiarthritic; antiviral;
KW antiinflammatory; antithyroid; antiallergic; antibacterial; cardiac;
KW dermatological; neuroprotective; thrombolytic; coagulant; neotropic;
KW vasotropic; antipapillary; antidiabetic; antirheumatic; antithyroid;
KW immune disorder; haematopoietic cell disorder; autoimmune disorder;
KW allergic reaction; graft versus host disease; organ rejection;
KW haemostatic; thrombolytic; cardiovascular disorder; infection;
KW neurological disease; drug screening; ss.
XX
OS Homo sapiens.
XX
PN WO200055350-A1.
XX
PD 21-SEP-2000.
XX
PF 08-MAR-2000; 2000WO-US05882.
XX
PR 12-MAR-1999; 99US-0124270.
XX
PA (HUMA-) HUMAN GENOME SCI INC.
XX
PI Rosen CA, Ruben SM;
XX
DR WPI; 2000-587533/55.
XX
DR P-PSDB; AAB43428.
XX
PT Novel isolated nucleic acids comprising sequences encoding peptides
PT useful for treating or diagnosing e.g. cancer -
XX
PS Claim 1; Page 633-634; 2352pp; English.
XX
CC AAC77607 to AAC78448 encode the human cancer associated proteins given
CC in AAB43398 to AAB44239. The proteins can have activities based on the
CC tissues and cells the genes are expressed in. Example of activities
CC include: cytostatic; proliferative; vulnery; immunomodulator;
CC antidiabetic; antiasthmatic; antirheumatic; antiarthritic;
CC antiinflammatory; antithyroid; antiallergic; antibacterial; cardiac;
CC dermatological; neuroprotective; thrombolytic; coagulant;
CC neotropic; vasotropic; antipapillary and angiogenic. The
CC polynucleotides and polypeptides can be used for preventing, treating or
CC ameliorating medical conditions and diagnosing pathological conditions.
CC Polynucleotides, polypeptides, antibodies, agonists and antagonists from
CC the present invention may be used to treat immune disorders by activating
CC or inhibiting the proliferation, differentiation or mobilisation of
CC immune cells; to treat disorders of haematopoietic cells, autoimmune
CC disorders, allergic reactions, graft versus host disease and organ
CC rejection, modulate haemostatic or thrombolytic activity, modulate
CC inflammation, cancers, cardiovascular disorders, neurological disease and
CC bacterial or viral infections. The peptides, nucleotides, antibodies,
CC agonists and antagonists may be also be used in drug screens. AAC78449 to
CC AAC78457 and AAB44240 represent sequences used in the exemplification of
CC the present invention.
XX
SQ Sequence 2059 BP; 505 A; 548 C; 552 G; 451 T; 3 other;
XX
Query Match 4.2%; Score 41.8; DB 21; Length 2059;
Best Local Similarity 61.5%; Pred. No. 2.2;
Matches 67; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
XX 884 CAGGCCAGGAGCTCAGGCCATCAGCGTGCATGTTCTGTTGAGTTCATGCACAA 943

Db 1945 CAGGCCAGGAGCTCAGGCCATCAGCGTGCATGTTCTGTTGAGTTCATGCACAA 2004
QY 944 GAGTGTGTTTGTAGCTGCTTGTCCACAAAAAAGAAAAAAGAAAAAAGAAAAA 992
Db 2005 TAAAGGCTGAGCTCTTATCTTGCACAAAAAAGAAAAAAGAAAAAAGAAAAA 2053

RESULT 34
ABX63699/c
ID ABX63699 standard; cDNA; 2672 BP.
XX
AC ABX63699;
XX
DT 26-FEB-2003 (first entry)
XX
DE Human cDNA #699 differentially expressed in activated vascular tissue.
XX
KW Human; gene; ss; vascular tissue; cytostatic; atherosclerosis;
KW cardiac; hypotensive; antidiabetic; gynaecological; vasotropic;
KW cerebroprotective; gene therapy; vascular disease; cancer; coronary;
KW artery disease; hypertension; diabetes; pre-eclampsia; restenosis;
KW ischaemia-reperfusion injury; stroke;
XX
OS Homo sapiens.
XX
PN US2002137081-A1.
XX
PD 26-SEP-2002.
XX
PF 08-JAN-2002; 2002US-0044090.
XX
PR 28-JUL-2000; 2000US-222469P.
XX
PR 08-JAN-2001; 2001US-260483P.
XX
PA (BAND/) BANDMAN O.
XX
PI Bandman O;
XX
DR WPI; 2003-110597/10.
XX
PT Combination for diagnosing, staging, treating, or monitoring the
PT progression of treatment of a vascular disease, e.g. atherosclerosis,
PT comprises several cDNAs that are differentially expressed in activated
PT vascular tissue -
XX
PS Claim 1; Page -; 18pp; English.
XX
CC This invention relates to a combination comprising several cDNAs that
CC are differentially expressed in activated vascular tissue. The invention
CC also discloses a high throughput method for detecting differentially
CC expressed cDNAs in a sample. The cDNAs of the invention may have
CC antiatherosclerotic; cytostatic; cardiac; hypotensive; antidiabetic;
CC gynaecological; vasotropic and cerebroprotective activities and may be
CC used in gene therapy. The cDNAs of the invention may be used in a
CC high-throughput methods for detecting differential expression of one or
CC more cDNAs in a sample, or screening several molecules or compounds to
CC identify a molecule or compound that specifically binds a cDNA of the
CC invention. A protein encoded by the cDNA may be used to screen several
CC molecules or compounds to identify a ligand that specifically binds to
CC the protein, or to produce or purify an antibody to the protein that can
CC be used to detect a protein in a sample or purify a natural or
CC recombinant protein from a sample. The nucleotides may be useful for
CC diagnosing, staging, treating, or monitoring the progression of
CC treatment of a vascular disease, e.g. atherosclerosis, cancer, coronary
CC artery disease, hypertension, diabetes, pre-eclampsia, ischaemia-
CC reperfusion injury, restenosis, or stroke. The cDNAs can also be used
CC for large-scale genetic or gene expression analysis of several new
CC nucleic acid molecules. Antibodies to the proteins encoded by the
CC cDNAs are useful for diagnosing pre-pathologic disorders, and chronic
CC or acute diseases associated with abnormalities in the expression,
CC amount or distribution of the protein. The present sequence
CC represents a cDNA of the invention that is differentially expressed in

activated vascular tissue.

Note: The sequence data for this patent did not form part of the specification, but was obtained in electronic format directly from USPTO at <http://seqdata.uspto.gov/sequence.html?DocID=20020137081>.

Sequence 2672 BP; 432 A; 880 C; 907 G; 453 T; 0 other;

Query Match 4.2%; Score 41.8; DB 25; Length 2672;
 Best Local Similarity 51.3%; Pred. No. 2.4;
 Matches 97; Conservative 0; Mismatches 92; Indels 0; Gaps 0;

QY 410 GATGCACTGGAGCTGTGGTCCAGCCAGTGTCTGAGTGTGGGGCCCAACTTCACTCTGCA 469
 DB 2220 GCTGGGCTGGCAGATGCAGTCCACGGCGCTGTGCTAGCCAGGACACGTGGCCAGGACC 2161

QY 470 GGACAGAGGGGCGAGCCCGCCAGCCGAGACCCGGCGGCTGTGCTAGCCAGGACACGTGGCCAGGACC 529
 DB 2160 CCGCAGAGGGGCGCCCGCAGACCCGGCGGCTGTGCTAGCCAGGACACGTGGCCAGGACC 2101

QY 530 TATCACCACAGCCTGTATGGGAGGATGGGCGAGTGGCGAGTCCACCTGCAGCAGAGACCATGCA 589
 DB 2100 TGCCACGAAACCGCCTTATGCCCATCGCGCTGGGGGGCGAGTAGCGTCTGCACCGA 2041

QY 590 CAGGCAGCC 598
 DB 2040 CAGGGAGGC 2032

RESULT 35
 AAHL1559/c
 ID AAHL1559 standard; cDNA; 3471 BP.
 AC AAHL1559;
 DT 26-JUN-2001 (first entry)
 XX Human cDNA sequence SEQ ID NO:13847.
 DE Human; primer; detection; diagnosis; antisense therapy; gene therapy; ss.
 KW Homo sapiens.
 OS
 EN EP1074617-A2.
 XX
 PD 07-FEB-2001.
 XX
 PF 28-JUL-2000; 2000EP-0116126.
 XX
 PR 29-JUL-1999; 99JP-0248036.
 PR 27-AUG-1999; 99JP-0300253.
 PR 11-JAN-2000; 2000JP-0118776.
 PR 02-MAY-2000; 2000JP-0183767.
 PR 09-JUN-2000; 2000JP-0241899.
 XX
 PA (HELIX-) HELIX RES INST.
 XX
 PI Ota T, Isogai T, Nishikawa T, Hayashi K, Saito K, Yamamoto J;
 PI Ishii S, Sugiyama T, Wakamatsu A, Nagai K, Otsuki T;
 XX
 DR WPI; 2001-318749/34.
 XX
 PT Primer sets for synthesizing polynucleotides, particularly the 5602
 PT full-length cDNAs defined in the specification, and for the detection
 PT and/or diagnosis of the abnormality of the proteins encoded by the
 PT full-length cDNAs -
 XX
 ES Claim 8; SEQ ID 13847; 2537bp + CD ROM; English.
 XX
 CC The present invention describes primer sets for synthesizing 5602
 CC full-length cDNAs defined in the specification. Where a primer set
 CC comprises: (a) an oligo-dT primer and an oligonucleotide complementary
 CC to the complementary strand of a polynucleotide which comprises one of
 CC the 5602 nucleotide sequences defined in the specification, where the

oligonucleotide comprises at least 15 nucleotides; or (b) a combination
 of an oligonucleotide comprising a sequence complementary to the
 complementary strand of a polynucleotide which comprises a 5'-end
 sequence and an oligonucleotide comprising a sequence complementary to a
 polynucleotide which comprises a 3'-end sequence, where the
 oligonucleotide comprises at least 15 nucleotides and the combination of
 the 5'-end sequence/3'-end sequence is selected from those defined in
 the specification. The primer sets can be used in antisense therapy and
 in gene therapy. The primers are useful for synthesizing polynucleotides,
 particularly full-length cDNAs. The primers are also useful for the
 detection and/or diagnosis of the abnormality of the proteins encoded by
 the full-length cDNAs. The primers allow obtaining of the full-length
 cDNAs easily without any specialised methods. AAH03166 to AAH13628 and
 AAH13633 to AAH18742 represent human cDNA sequences; AAH92446 to
 AAH95893 represent human amino acid sequences; and AAH13629 to AAH13632
 represent oligonucleotides, all of which are used in the exemplification
 of the present invention.

Sequence 3471 BP; 568 A; 1088 C; 1233 G; 582 T; 0 other;

Query Match 4.2%; Score 41.8; DB 22; Length 3471;
 Best Local Similarity 51.3%; Pred. No. 2.6;
 Matches 97; Conservative 0; Mismatches 92; Indels 0; Gaps 0;

QY 410 GATGCACTGGAGCTGTGGTCCAGCCAGTGTCTGAGTGTGGGGCCCAACTTCACTCTGCA 469
 DB 3027 GCTGGGCTGGCAGATGCAGTCCACGGCGCTGTGCTAGCCAGGACACGTGGCCAGGACC 2968

QY 470 GGACAGAGGGGCGAGCCCGCCAGCCGAGACCCGGCGGCTGTGCTAGCCAGGACACGTGGCCAGGACC 529
 DB 2967 CCGCAGAGGGGCGCCCGCAGACCCGGCGGCTGTGCTAGCCAGGACACGTGGCCAGGACC 2908

QY 530 TATCACCACAGCCTGTATGGGAGGATGGGCGAGTGGCGAGTCCACCTGCAGCAGAGACCATGCA 589
 DB 2907 TGCCACGAAACCGCCTTATGGCCATCGCGCTGGGGGGCGAGTAGCGTCTGCACCGA 2848

QY 590 CAGGCAGCC 598
 DB 2847 CAGGGAGGC 2839

RESULT 36
 ABL53715/c
 ID ABL53715 standard; cDNA; 3541 BP.
 AC ABL53715;
 XX
 DT 24-JUN-2002 (first entry)
 XX Human poly(ADP-ribose) polymerase p150 cDNA.
 DE Human poly(ADP-ribose) polymerase; PARP; p150; human; enzyme; diabetes;
 KW cerebral ischaemia; Alzheimer's disease; Parkinson's disease;
 KW systemic lupus erythematosus; arthritis; tumour; therapy;
 KW diagnosis; cytostatic; vasotropic; antidiabetic; neurotropic;
 KW neuroprotective; antiparkinsonian; antiarthritic; dermatological;
 KW immunosuppressive; antiinflammatory; expressed sequence tag; EST;
 KW gene; chromosome 8; ss.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT CDS 122..3199
 FT /*tag= a
 FT /product= "Poly(ADP-ribose) polymerase"
 XX
 XX WO200222792-A2.
 XX
 PD 21-MAR-2002.
 XX
 PF 11-SEP-2001; 2001WO-EPI0494.
 XX
 PR 12-SEP-2000; 2000EP-0119849.


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RESULT 38
ABN97975
XX ABN97975 standard; DNA; 44100 BP.
XX
XX AC ABN97975;
XX
XX DT 01-AUG-2002 (first entry)
XX
XX DE Human retroviral sequence RAM75.
XX
XX KW Autoimmune disease; HERV-7q; chromosome 7q; immunotherapy;
XX multiple sclerosis; ds.
XX
XX OS Human retrovirus.
XX
XX PN WO9967395-A1.
XX
XX PD 29-DEC-1999.
XX
XX PF 23-JUN-1999; 99WO-FR01513.
XX
XX PR 23-JUN-1998; 98FR-0007920.
XX
XX (INERM ) INSERM INST NAT SANTE & RECH MEDICALE.
XX
XX PI Alliel PM, Perin J, Rieger F;
XX
XX DR WPI; 2000-160587/14.
XX
XX PT New nucleic acid sequences of human endogenous retrovirus, HERV-7q,
XX used for diagnosis, treatment and prevention of autoimmune and
XX neurological diseases -
XX
XX PS Claim 15; Fig 14; 225pp; French.
XX
XX CC The present invention relates to new nucleic acid sequences of human
XX endogenous retrovirus, HERV-7q, which is located on chromosome 7q.
XX Regulatory elements associated with HERV-7q may alter expression of other
XX genes (even remote genes) on the same chromosome, inducing immunological
XX and/or neurological changes (which may be pathological or protective/
XX curative). HERV-7q peptides can be used to improve efficiency of the
XX immune response, e.g. in immunotherapy. HERV-7q peptides and their coding
XX sequences can be used in immunogenic or vaccinating compositions, for
XX protection against autoimmune diseases, particularly multiple sclerosis.
XX The peptides may also be used (by sequence comparison) to detect/identify
XX endogenous retroviruses that are abnormally expressed in cancer,
XX neuropathologies or other autoimmune diseases. The present sequence was
XX used to illustrate the invention.
XX
XX SQ Sequence 44100 BP; 13051 A; 8074 C; 8534 G; 14441 T; 0 other;

Query Match 4.0%; Score 39.6; DB 21; Length 44100;
Best Local Similarity 77.4%; Pred. No. 21;
Matches 48; Conservative 0; Mismatches 14; Indels 0; Gaps 0;

QY 920 TCGTATTGGAGTTCATGCATAATGAGTGTGTTTGGCTTGGCCACAAAAA 979
DB 10427 TTGTATGGGATTCATGTTAAATGAGTAGATTTTAACTACTCTTACCACAAAACAA 10486
QY 980 AA 981
DB 10487 AA 10488

RESULT 39
ABN22420
XX ABN22420 standard; cDNA; 276 BP.
XX
XX AC ABN22420;
XX
XX DT 24-JUN-2002 (first entry)
XX

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XX DE Human ORFX polynucleotide sequence SEQ ID NO:13317.
XX
XX KW Human; open reading frame; ORFX; gene therapy; cancer; cirrhosis;
XX hyperproliferative disorder; psoriasis; benign tumour; haemorrhage;
XX degenerative disorder; osteoarthritis; neurodegenerative disorder;
XX cardiovascular disease; diabetes mellitus; systemic lupus erythematosus;
XX hypertension; hypothyroidism; cholesterol ester storage disease;
XX immune deficiency; immune disorder; infectious disease;
XX autoimmune disorder; rheumatoid arthritis; autoimmune thyroiditis;
XX myasthenia gravis; gene; ss.
XX
XX OS Homo sapiens.
XX
XX PN WO200192523-A2.
XX
XX PD 06-DEC-2001.
XX
XX PF 29-MAY-2001; 2001WO-US10836.
XX
XX PR 30-MAY-2000; 2000US-206132P.
XX
XX PR 29-AUG-2000; 2000US-228716P.
XX
XX PA (CURA-) CURAGEN CORP.
XX
XX XX Shinkets RA, Leach MD;
XX
XX PI WPI; 2002-106308/14.
XX
XX DR P-PSDB; ABP06668.
XX
XX PT Novel human polypeptides and polynucleotides useful for diagnosing,
XX preventing and treating cardiovascular disease, neurodegenerative,
XX hyperproliferative disorders and autoimmune disorders -
XX
XX PS Disclosure; SEQ ID 13317; 1037pp; English.
XX
XX CC The present invention describes substantially purified human proteins
XX (referred to as open reading frame, ORFX, where X is 1-11491 (see Table 1
XX in the specification). ABN15762 to ABN27252 encode the human ORFX
XX proteins given in ABP00010 to ABP11500. ORFX proteins are useful for
XX treating or preventing a pathology associated with an ORFX-associated
XX disorder in humans, and in the manufacture of a medicament for treating a
XX syndrome associated with ORFX-associated disorder. ORFX polynucleotide
XX sequences can be used in gene therapy. ORFX sequences can be used in the
XX treatment of cancer, hyperproliferative disorders, cirrhosis of liver,
XX psoriasis, benign tumours, keloid, degenerative disorders, haemorrhage,
XX osteoarthritis, neurodegenerative disorders, disorders related to organ
XX transplantation, cardiovascular diseases, diabetes mellitus, systemic
XX lupus erythematosus, hypertension, hypothyroidism, cholesterol ester
XX storage disease, various immune deficiencies and disorders, infectious
XX diseases, autoimmune disorders such as multiple sclerosis, rheumatoid
XX arthritis, autoimmune thyroiditis, myasthenia gravis, graft-versus-host
XX disease and autoimmune inflammatory eye disease. ORFX proteins are also
XX useful for treating burns, incisions, ulcers, for treating osteoporosis,
XX bone degenerative disorders, or periodontal disease, and for gut
XX protection or regeneration and treatment of lung or liver fibrosis,
XX reperfusion injury in various tissues and conditions resulting from
XX systemic cytokine damage.
XX
XX N.B. The sequence data for this patent did not form part of the printed
XX specification, but was obtained in electronic format directly from WIPO
XX at ftp.wipo.int/pub/published_pct_sequences.
XX
XX SQ Sequence 276 BP; 43 A; 105 C; 86 G; 40 T; 2 other;

Query Match 4.0%; Score 39.4; DB 24; Length 276;
Best Local Similarity 50.8%; Pred. No. 4.4;
Matches 94; Conservative 0; Mismatches 91; Indels 0; Gaps 0;

QY 338 TCCAGACCTGCTCACCTACTTCTGCGGGCGTCTCTCCACCTCAGTCCCATGTGGACAG 397
DB 9 TCCCCACCTGCTCGAGCGGCTCGACCTCGGCCACACCACTCGCGCAACCGCGTGATCAT 68
QY 398 TGCAGGCTACAGATGCACCTGGGAGCTGTGTGTCACGCCAGTGTCTGAGCTGGGGCAA 457

```

Db 69 GGGCTCGATGCACACCGCATGGAGGATCGTCCAGCACCTGCCGAGCTGGCGGCTA 128
 QY 458 CTTCACTCTGCAGACAGAGGGGCGAGGCCCGCCAGGCTGGAGATGATGTCGCCAGGCGTCTC 517
 Db 129 CTTCCGCGAGCGGGCCCAAGGCTGGCGCTGTCCGCTCAACCGCGCGCTACCGCCCAAC 188
 QY 518 GGGCA 522
 Db 189 TGGCA 193
 RESULT 40
 AAX80672
 ID AAX80672 standard; cDNA; 3285 BP.
 XX AC AAX80672;
 XX DT 22-OCT-1999 (first entry)
 XX DE Clone nm103_4 encoding secreted protein-nm103_4.
 XX KW Secreted protein; cytokine; cell proliferation; immune stimulation;
 KW vaccine; immune suppression; haematopoiesis; tissue growth; activin;
 KW inhibin; chemotaxis; chemokinesis; haemostasis; thrombolytic; ss;
 KW receptor; ligand; anti-inflammatory; cadherin; tumour; gene therapy.
 OS Homo sapiens.
 XX FH Key Location/Qualifiers
 FT CDS 2294..2848
 FT /*tag= a
 FT /product= "Secreted protein nm103_4"
 XX PN W09928335-A1.
 XX PD 10-JUN-1999.
 XX PF 02-DEC-1998; 98WO-US25512.
 XX PR 30-NOV-1998; 98US-0203106.
 XX PR 04-DEC-1997; 97US-0067454.
 XX PA (GEMY) GENETICS INST INC.
 XX PI Agostino MJ, Clark HF, Collins-Racie LA, Evans C;
 PI Fechtel K, Jacobs K, Lavallie ER, McCoy JM, Merberg D;
 PI Steininger RJ, Treacy M, Wong GG;
 XX WPI; 1999-385352/32.
 XX P-PSDB; AAV26037.
 XX PT New polynucleotides encoding secreted human proteins
 XX PS Claim 28; Pages 111-112; 124pp; English.
 XX CC The present sequence is a known clone nm103_4 (deposited as ATCC
 CC 98600) isolated from human foetal kidney cDNA library. It encodes
 CC secreted protein nm103_4. Recombinant secreted proteins can be produced
 CC by transforming host cells and culturing them under suitable conditions.
 CC The polynucleotide and protein are predicted to have biological
 CC activities which would make them suitable for treating, preventing or
 CC ameliorating medical conditions in humans and animals. Some predicted
 CC biological activities include cytokine and cell proliferation/
 CC differentiation activity, immune stimulating (e.g. as vaccines) or
 CC suppressing activity, haematopoiesis regulating activity, tissue growth
 CC activity, activin/inhibin activity, chemotactic/chemokinetic activity,
 CC hemostatic and thrombolytic activity, receptor/ligand activity, anti-
 CC inflammatory activity, cadherin/tumour invasion suppressor activity, and
 CC tumour inhibition activity. The polynucleotide can be used for gene
 CC therapy.
 XX CC Sequence 3285 BP; 756 A; 813 C; 705 G; 1011 T; 0 other;
 SQ

Query Match 4.0%; Score 39.4; DB 20; Length 3285;
 Best Local Similarity 72.5%; Pred. No. 10;
 Matches 66; Conservative 0; Mismatches 21; Indels 4; Gaps 1;
 QY 891 GAGGACTGCGAGGCATCAGCGTCGACCTGTTGATTTGGAGTTTCATGCAAAATGAGTGTG 950
 Db 3049 GAGGAATGTAGGTAATAAATTCGACTGTT-----TTGGGAGTTCTCTAATGACTAGA 3104
 QY 951 TTTTAGCTGCTCTTGGCCACCAAAAAA 981
 Db 3105 CTTCACTGCTCTTGGCACAATACTTAAA 3135
 RESULT 41
 AAS59222
 ID AAS59222 standard; cDNA; 3285 BP.
 XX AC AAS59222;
 XX DT 16-JAN-2002 (first entry)
 XX DE Human cDNA encoding a secreted protein nm103_4.
 XX KW Human; secreted protein; ss; antiinflammatory; immunosuppressive;
 KW neotropic; neuroprotective; antiarthritic; antimicrobial; vulnerary;
 KW cyostatic; antidiabetic; virucide; antiinfertility; anticonvulsant;
 KW vasotropic; antiparkinsonian; immunostimulant; dermatological;
 KW antirheumatic; antitumor; antiulcer; osteopathic; tranquiliser;
 KW cerebroprotective; cytokine; cell proliferation; cell differentiation;
 KW immune deficiency; severe combined immunodeficiency; SCID; tumour;
 KW autoimmune disorder; multiple sclerosis; rheumatoid arthritis;
 KW graft-versus-host disease; myeloid deficiency; wound healing; ulcer;
 KW periodontal disease; osteoporosis; osteoarthritis; Alzheimer's disease;
 KW Parkinson's disease; Huntington's disease; infection; cardiac disease;
 KW stroke; sepsis; inflammatory bowel disease; contraceptive; immunogen;
 KW food supplement.
 XX OS Homo sapiens.
 XX WO200175068-A2.
 XX 11-OCT-2001.
 XX PF 22-MAR-2001; 2001WO-US09369.
 XX PR 30-MAR-2000; 2000US-0539330.
 XX PR 04-DEC-2000; 2000US-0729674.
 XX PA (GEMY) GENETICS INST INC.
 XX PI Jacobs K, McCoy JM, Lavallie E, Collins-racie LA, Evans C;
 PI Treacy M, Agostino MJ, Steininger RJ, Spaulding V, Wong GG;
 PI Clark H, Fechtel K, Merberg D;
 XX WPI; 2001-639363/73.
 XX P-PSDB; AAU39004.
 XX PT Secreted human proteins, useful as vaccine for treating various
 XX diseases such as autoimmune disorders (e.g. multiple sclerosis), and
 XX nervous system disorders (e.g. stroke) -
 XX PS Disclosure; Page 476-477; 619pp; English.
 XX CC The invention relates to novel human secreted proteins, the nucleic
 CC acids encoding them. The protein may exhibit cytokine, cell proliferation
 CC or cell differentiation activity or may induce production of other
 CC cytokines in certain cell populations and may exhibit immune stimulating
 CC or immune suppressing activity, which is useful for the treatment of
 CC various immune deficiencies and disorders e.g. severe combined
 CC immunodeficiency (SCID), autoimmune disorders e.g. multiple sclerosis,
 CC systemic lupus erythematosus, rheumatoid arthritis, autoimmune pulmonary
 CC inflammation. The proteins are also useful in the treatment of diseases


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XX DT 09-APR-2001 (first entry)
XX DE Novel human polynucleotide, SEQ ID NO: 788.
XX KW Human; cytostatic; gene therapy; colon cancer; prostate cancer;
XX KW breast cancer; lung cancer; cancer detection; ss.
XX OS Homo sapiens.
XX PN WO200102568-A2.
XX PD 11-JAN-2001.
XX PF 30-JUN-2000; 2000WO-US18374.
XX PR 02-JUL-1999; 99US-0142310.
XX PR 02-JUL-1999; 99US-0142311.
XX PA (CHIR ) CHIRON CORP.
XX PA (HYSE-) HYSEQ INC.
XX PI Williams LT, Escobedo J, Innis MA, Garcia PD, Klinger J, Kassam A;
XX PI Reinhard C, Randazzo F, Kennedy GC, Pot D, Lamson G, Drmanac R;
XX PI Ckerjakov R, Drmanac S, Dickson M, Labat I, Leshkowitz D;
XX PI Kita D, Garcia V, Jones LM, Strache-Crain B;
XX DR WPI; 2001-091805/10.
XX KW Library of polynucleotides for diagnosing a cancerous state of a
XX PT mammalian cell and detecting cancer, particularly of the colon or
XX PT prostate, comprises 3351 human polynucleotide sequences -
XX PS Claim 9; Page 657-658; 1046pp; English.
XX CC The present sequence is one of 3351 sequences in a library of human
XX CC polynucleotides. The library is used to detect differentially expressed
XX CC genes correlated with a cancerous state of a mammalian cell and can
XX CC detect colon, prostate, breast and lung cancer. The library can be used
XX CC to produce probes for detection of mRNA and to produce additional copies
XX CC of the polynucleotides. The probes can be used for chromosome mapping of
XX CC the polynucleotide and for detection of transcription levels. Ribozymes
XX CC or antisense oligonucleotides can be generated. The polynucleotides and
XX CC their gene products are used as genetic or biochemical markers (e.g. in
XX CC blood or tissues) that will detect the earliest changes along the
XX CC carcinogenesis pathway and/or monitor the efficacy of therapies and
XX CC preventive interventions. The polynucleotides, polypeptides and
XX CC antibodies against them can be used in pharmaceutical compositions to
XX CC treat the cancers and proliferative disorders such as neoplasia,
XX CC dysplasia and hyperplasia.
XX SQ Sequence 394 BP; 103 A; 90 C; 115 G; 86 T; 0 other;

Query Match 4.0%; Score 39.2; DB 22; Length 394;
Best Local Similarity 66.7%; Pred. No. 5.5;
Matches 56; Conservative 0; Mismatches 28; Indels 0; Gaps 0;

QY 909 GCGTGCACCTGTCGTATTTGAGTTTCATGCAAAATGAGTGTGTTTATGCTGCTTGCCTA 968
Db 393 GCGCGCGCCGCTCTGTGGTGAGACATACACACACAGAGTGAGTGTGTTTCCCTC 334

QY 969 CAAAAAATAAAAAAAAAAAAAA 992
Db 333 CACAAAAAATAAAAAAAAAAAGAA 310

RESULT 44
ID AAC98202
AC AAC98202 standard; cDNA; 1639 BP.
AC AAC98202;
XX DT 09-MAR-2001 (first entry)

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XX XX Human colon cancer antigen nucleotide sequence SEQ ID NO:212.
XX DE Human; colon cancer; colon cancer antigen; diagnosis; detection;
XX KW identification; cytostatic; cardioactive; neuroprotective; vulnerary;
XX KW immunomodulatory; muscular; gynaecological; gastrointestinal;
XX KW nephrotropic; antiinfective; antibacterial; gene therapy; wound;
XX KW neural disorder; immune system disorder; muscular disorder;
XX KW reproductive disorder; gastrointestinal disorder; renal disorder;
XX KW infectious disease; cardiovascular disorder; ss.
XX OS Homo sapiens.
XX PN WO200055351-A1.
XX PD 21-SEP-2000.
XX PF 08-MAR-2000; 2000WO-US05883.
XX PR 12-MAR-1999; 99US-0124270.
XX PA (HUMA-) HUMAN GENOME SCI INC.
XX PI Rosen CA, Ruben SM;
XX PR WPI; 2000-587534/55.
XX DR P-PSDB; AAB53445.
XX KW Colon cancer associated gene sequences, referred to as colon cancer
XX PT antigens, useful for the treatment, prevention, and diagnosis of colon
XX PT disorders such as colon cancer -
XX PS Claim 1; Page 636; 2104pp; English.
XX CC AAC97991 to AAC98763 encode the human colon cancer associated proteins,
XX CC called human colon cancer antigens, given in AAB53234 to AAB54006. The
XX CC human colon cancer antigens can have cytostatic, cardioactive, muscular;
XX CC neuroprotective, immunomodulatory, gynaecological, gastrointestinal,
XX CC vulnerary, nephrotropic, antiinfective and antibacterial activities, and
XX CC can be used in gene therapy. The colon cancer antigen polynucleotides,
XX CC proteins and antibodies to the proteins are useful for the prevention,
XX CC treatment and diagnosis of colon disorders, such as colon cancer. The
XX CC polynucleotides may be used in diagnostics and research, such as for
XX CC chromosome identification, and as hybridisation probes. The proteins
XX CC may also be used to prevent diseases such as neural disorders, immune
XX CC system disorders, muscular disorders, reproductive disorders,
XX CC gastrointestinal disorders, wounds, renal disorders, infectious
XX CC diseases, and cardiovascular disorders. AAC98764 to AAC98772 and
XX CC AAB54007 represent sequences used in the exemplification of the present
XX CC invention.
XX SQ Sequence 1639 BP; 410 A; 398 C; 357 G; 469 T; 5 other;

Query Match 3.9%; Score 38.8; DB 21; Length 1639;
Best Local Similarity 64.4%; Pred. No. 11;
Matches 58; Conservative 0; Mismatches 32; Indels 0; Gaps 0;

QY 903 CCATCAGCGTGCACCTGTCGTATTTGAGTTTCATGCAAAATGAGTGTGTTTATGCTGCTC 962
Db 1501 CCAGCACCACTTATGAGTCTCAAGTTTATTTATTTGCAATAAAAGTCTTTATGCCGCTT 1560

QY 963 TTGCCCAAAAAAAAAAAAAAAAAAAAAA 992
Db 1561 TTCTCAAAAAAAAAAAAAAAAAAAAAA 1590

RESULT 45
ID AABK33656
AC AABK33656 standard; cDNA; 2025 BP.
AC AABK33656;
XX DT 08-MAY-2002 (first entry)

```

XX DE cDNA encoding human PRO protein, Seq ID No 241.
 XX KW Human; secreted protein; PRO; tumour; lung cancer; colon cancer;
 XX KW breast cancer; prostate tumour; rectal tumour; liver tumour;
 KW KW pericyte cell proliferation; chondrocyte cell proliferation;
 KW KW tumour necrosis factor-alpha; gene; ss.
 XX OS Homo sapiens.
 XX KW WO200208288-A2.
 XX PD 31-JAN-2002.
 XX XX 29-JUN-2001; 2001WO-US21066.
 XX PF 20-JUL-2000; 2000US-219556P.
 XX PR 25-JUL-2000; 2000US-220585P.
 XX PR 25-JUL-2000; 2000US-220605P.
 XX PR 25-JUL-2000; 2000US-220607P.
 XX PR 25-JUL-2000; 2000US-220624P.
 XX PR 25-JUL-2000; 2000US-220638P.
 XX PR 25-JUL-2000; 2000US-220664P.
 XX PR 25-JUL-2000; 2000US-220666P.
 XX PR 26-JUL-2000; 2000US-220893P.
 XX PR 28-JUL-2000; 2000WO-US20710.
 XX PR 23-AUG-2000; 2000WO-US23522.
 XX PR 24-AUG-2000; 2000WO-US23328.
 XX PR 15-SEP-2000; 2000US-000000P.
 XX PR 10-NOV-2000; 2000WO-US30873.
 XX PR 28-NOV-2000; 2000US-253646P.
 XX PR 01-DEC-2000; 2000WO-US32678.
 XX PR 20-DEC-2000; 2000US-0747259.
 XX PR 20-DEC-2000; 2000WO-US34956.
 XX PR 28-FEB-2001; 2001WO-US06520.
 XX PR 10-MAY-2001; 2001US-0854280.
 XX PR 25-MAY-2001; 2001WO-US17092.
 XX PA (GETH) GENENTECH INC.
 XX XX Baker KP, Desnoyers L, Gerritsen ME, Goddard A, Godowski PJ;
 PI PI Grimaldi JC, Gurney AL, Smith V, Stephan JF, Watanabe CK, Wood WI;
 XX P-PSDB; AAU83712.
 XX WPI; 2002-172001/22.
 XX DR One hundred and twenty two nucleic acids encoding PRO polypeptides,
 XX DR useful for treating a PRO related disorder and for diagnosing tumours
 XX PT such as lung cancer, colon cancer, breast tumour, prostate tumour, rectal
 XX PT tumour or liver tumour -
 XX PS Claim 2; Figure 241; 359pp; English.
 XX XX The invention relates to one hundred and twenty two nucleic acids
 CC encoding PRO polypeptides. The sequences of the 122 PRO polynucleotides
 CC encode human secreted proteins. The PRO nucleic acids, polypeptides,
 CC agonists and antagonists are useful for treating a PRO related disorder.
 CC The PRO polypeptides are useful for diagnosing tumours, especially lung
 CC cancer, colon cancer, breast tumour, prostate tumour, rectal tumour or
 CC liver tumour. The PRO polypeptides are useful for stimulating the
 CC proliferation of, or gene expression, in pericyte cells, for stimulating
 CC the proliferation or differentiation of chondrocyte cells, for
 CC stimulating the release of tumour necrosis factor-alpha from human blood,
 CC for stimulating or inhibiting the proliferation of normal human dermal
 CC fibroblast cells. The PRO polypeptide may also be used as molecular
 CC weight markers and for tissue typing. The PRO nucleic acids have
 CC applications in molecular biology, including use as hybridisation probes,
 CC and in chromosome and gene mapping. ABK33536-ABK33657 represent human
 CC PRO protein coding sequences of the invention.
 XX SQ Sequence 2025 BP; 471 A; 592 C; 584 G; 378 T; 0 other;
 XX Query Match 3.9%; Score 38.8; DB 24; Length 2025;

Best Local Similarity 60.4%; Pred. No. 12;
 Matches 64; Conservative 0; Mismatches 42; Indels 0; Gaps 0;
 QY 887 GCGAGAGACTGCGAGCCATCAGCTGCTTCTGATTTTCGAGTTTCATGCAAAATGAG 946
 DB 1886 GGTGCTGGCCAGCGGACTCAGGGCGGAGGCTTCCCGGGGAGCAGGAGAAATAA 1945
 QY 947 TGTGTTTGTAGTCTTTCGCCACAAAAAATAAAAAAAAAAAAAA 992
 DB 1946 GGTCTTTGCTTTCTCCAGCAAAAAAATAAAAAAAAAAAAAA 1991

RESULT 46
 ABS69895/c
 ID ABS69895 standard; DNA; 56737 BP.
 XX AC ABS69895;
 XX DT 21-NOV-2002 (first entry)
 XX DE Human hypoxanthine-guanine phosphoribosyltransferase (HPRT) gene.
 XX KW Vector; adenovirus; adeno-associated; adenosine deaminase gene; receptor;
 KW adenosine deaminase deficiency; severe combined immune deficiency; PAH;
 KW beta-chain; haemoglobin gene; beta-thalassaemia; sickle cell disease;
 KW low density lipoprotein gene; familial hypercholesterolaemia;
 KW hypoxanthine-guanine phosphoribosyltransferase; Lesch-Nyhan syndrome;
 KW phenylalanine hydroxylase gene; gene therapy; phenylketonuria;
 KW dystrophin gene; muscular dystrophy; cystic fibrosis; immunostimulant;
 KW human cystic fibrosis transmembrane conductance regulator gene;
 KW antianemic; antilipemic; nootropic; cytostatic; dermatological;
 KW human; alpha-1-antitrypsin; lysosomal glucocerebrosidase; ADA; HPRT;
 KW lysosomal arylsulphatase A; omithine transcarbamylase; ARSA; OTC; NP;
 KW purin nucleoside phosphorylase; gene; ds.
 XX OS Homo sapiens.
 XX XX US2002102731-A1.
 XX PD 01-AUG-2002.
 XX XX 12-FEB-2001; 2001US-0782378.
 XX XX 02-OCT-2000; 2000US-237747P.
 XX PA (UNYNY) UNIV NEW YORK STATE RES FOUND.
 XX PI Hearing P, Bahou WF, Sandalon Z, Gnatenko DV;
 XX XX WPI; 2002-690619/74.
 XX PT Producing vector, by introducing vector having nucleotide sequence,
 PT adenovirus inverted terminal repeats and packaging sequence, and
 PT adeno-associated virus terminal repeat, into cell, and culturing cell
 XX -
 XX PS Disclosure; Page 93-117; 191pp; English.
 XX CC The present invention relates to a new method of producing a vector. The
 CC method involves introducing recombinant vector having nucleotide sequence
 CC (NS) having 5' and 3' end, left and right inverted terminal repeats of
 CC adenovirus flanking NS, adenovirus packaging sequence linked to inverted
 CC terminal repeat, and adeno-associated virus terminal repeat linked to 3'
 CC end of NS, into cell expressing adenovirus early gene lacking from
 CC vector; and culturing cell to produce another vector. The method is
 CC useful for generating vectors, especially MAD vectors. The method is
 CC useful in transferring nucleotide sequences of interest into a cell, for
 CC gene transfer applications (e.g. gene therapy) in vitro, ex vivo and in
 CC vivo. The nucleotide sequences are useful for treating diseases
 CC associated with it, i.e. adenosine deaminase gene associated with
 CC adenosine deaminase deficiency with severe combined immune deficiency,
 CC beta-chain of haemoglobin gene associated with beta-thalassaemia and
 CC sickle cell disease, receptor for low density lipoprotein gene associated

CC with familial hypercholesterolaemia, hypoxanthine-guanine
 CC phosphoribosyltransferase associated with Lesch-Nyhan syndrome,
 CC phenylalanine hydroxylase (PAH) gene associated with phenylketonuria,
 CC dystrophin gene associated with muscular dystrophy, and human cystic
 CC fibrosis transmembrane conductance regulator gene associated with cystic
 CC fibrosis. The present nucleic acid sequence represents a human disease
 CC gene sequence that was used in the methods of the invention.
 XX
 SQ Sequence 56737 BP; 15689 A; 11281 C; 11599 G; 18168 T; 0 other;
 Query Match 3.9%; Score 38.6; DB 24; Length 56737;
 Best Local Similarity 59.9%; Pred. No. 41;
 Matches 100; Conservative 0; Mismatches 64; Indels 3; Gaps 2;
 QY 819 GGGAGTGGGAGGCTCAGAGCAGCAAGCAGCAGCCATGTAGATGAACCGTCCAGAGAG 878
 Db 22684 GGGAGATGTAGCTCAGAGATCAAGTAGCAGATATGTAGATGAACAGTCTAGAAA 22625
 QY 879 CCAAGCAGGCGAGGAGCTCAGGCCATCAGCGTGCACCTTTCGTATTGGAGTTTCATGC 938
 Db 22624 TATAATGTACAACATGAAGTATATAGTATATAAAATTGTGCT-GTATTGGGATTCACAC 22566
 QY 939 AAAATGAGTGTGTTTAGCTGCTCTTCCACAAAAAAGAAAAA 985
 Db 22565 TAAATGA--GATTTAAGCTCCTCTTGCCACAAAAACAAAAA 22521
 RESULT 47
 ABQ77405
 ID ABQ77405 standard; DNA; 185695 BP.
 XX
 AC ABQ77405;
 XX
 DT 10-MAY-2003 (first entry)
 XX
 DE Human THBS1 DNA.
 XX
 KW Human; THBS1; vascular disease; cardiant; antiarteriosclerotic; stroke;
 KW cerebroprotective; gene therapy; coronary artery disease; ischaemia;
 KW myocardial infarction; peripheral vascular disease; pulmonary embolism;
 KW venous thromboembolism; forensic; paternity testing; GI12583762; gene;
 KW SNP; single nucleotide polymorphism; ds.
 XX
 OS Homo sapiens.
 XX
 FH Key Location/Qualifiers
 FT variation replace (53502,t)
 FT /*tag= a
 FT /standard name= "SNP"
 FT /note= "Single nucleotide polymorphism (ID G334u3)
 FT which does not change the THBS1 protein"
 XX
 PN WO2003016494-A2.
 XX
 PD 27-FEB-2003.
 XX
 PF 16-AUG-2002; 2002WO-US26343.
 XX
 PR 16-AUG-2001; 2001US-313097P.
 PR 05-OCT-2001; 2001US-327485P.
 PR 14-DEC-2001; 2001US-0020141.
 XX
 XX (VITI-) VITIVITY INC.
 XX
 PI McCarthy J, Ableson A;
 XX
 DR WPI; 2003-300617/29.
 DR P-PSDB; ABG74673.
 XX
 PT Identifying a subject as a candidate for a particular course of therapy
 PT to treat a vascular disease or disorder, e.g. stroke, myocardial
 PT infarction or ischemia by determining the identity of the nucleotide
 PT present at specific positions -

XX
 PS Claim 3; Fig 11; 568pp; English.
 XX
 CC This invention describes a novel method for identifying a subject as a
 CC candidate for a particular course of therapy to treat a vascular disease
 CC or disorder. The method comprises determining the identity of the
 CC nucleotide present at specific positions, or their complements, and
 CC identifying the subject as a candidate for a particular clinical course
 CC of therapy based on the identity of the nucleotide present in that
 CC specific position. The method can be used for identifying a subject who
 CC is a candidate for further diagnostic evaluation of a vascular disease or
 CC disorder and selecting a clinical course of therapy. The products of the
 CC invention have cardiant, antiarteriosclerotic and cerebroprotective
 CC activity and can be used for gene therapy. The methods disclosed are
 CC useful for treating a vascular disease, e.g. atherosclerosis, coronary
 CC artery disease, myocardial infarction, ischaemia, stroke, peripheral
 CC vascular diseases, venous thromboembolism and pulmonary embolism. The DNA
 CC sequences are useful as fingerprint for detecting different individuals
 CC within the same species applicable in forensic studies and paternity
 CC testing. This sequence encodes the human THBS1 gene represented in
 CC GI12583762, used to illustrate the method of the invention.
 XX
 SQ Sequence 185695 BP; 59388 A; 38636 C; 35664 G; 52007 T; 0 other;
 Query Match 3.9%; Score 38.4; DB 25; Length 185695;
 Best Local Similarity 53.7%; Pred. No. 68;
 Matches 101; Conservative 0; Mismatches 86; Indels 1; Gaps 1;
 QY 795 GAGGAGTTTGGGGGTTCCAGGATAGGGAGTGGGAGTCCAGAGCGCAAGCGCCAGCAGCC 854
 Db 112437 GATGTGTCGAGACTGTGTAGGAAATGGGGAGATATACATAAAGGATACAAAGTTGCAGAC 112496
 QY 855 ATGTAGATCAACCGTCCAGAGAGCCAGCAGCGCAGAGGACTGCGAGGCCATCAGCGTGC 914
 Db 112497 ATGTAAGATGAACAGTCTGGAGTCTTACATACACATGAGGACTATAGTTAGTAATAC 112556
 QY 915 ACTGTTTCGTATTGGAGTTCATGCAAAATGAGTGTGTTTACGTCTCTTGCACAAAAA 974
 Db 112557 TGCCTT-TTATTGAGATCTTGCACAAAAGAGTAGATTTTACGTGTCTTCCACACATA 112615
 QY 975 AAAAAAAA 982
 Db 112616 CGCAAGA 112623
 RESULT 48
 ABV61090/c
 ID ABV61090 standard; cDNA; 294 BP.
 XX
 AC ABV61090;
 XX
 DT 13-SEP-2002 (first entry)
 XX
 DE Human prostate expression marker cDNA 61081.
 XX
 KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
 KW pharmacogenomic marker; gene; ss.
 XX
 OS Homo sapiens.
 XX
 FN WO200160860-A2.
 XX
 XX 23-AUG-2001.
 XX
 PD 20-FEB-2001; 2001WO-US05171.
 XX
 PR 17-FEB-2000; 2000US-183319P.
 PR 16-MAR-2000; 2000US-189862P.
 PR 25-MAY-2000; 2000US-207454P.
 PR 09-JUN-2000; 2000US-211314P.
 PR 18-JUL-2000; 2000US-219007P.
 PR 13-DEC-2000; 2000US-255281P.
 XX

PA (MILL-) MILLENNIUM PREDICTIVE MEDICINE INC.
XX Schlegel R, Endege WO, Monahan JE;
XX WPI; 2001-662795/76.
XX Novel isolated nucleic acid molecule associated with cancerous state of
PT prostate cells and correlating with presence of prostate cancer, useful
PT for detecting presence of prostate cancer, stage of prostate cancer -
XX
XX Claim 1; Page 11605-11606; 11750pp; English.
XX
XX The invention relates to an isolated nucleic acid molecule (I) comprising
CC a nucleotide sequence given in Tables 1-9 (ABV00010-ABV62213) of the
CC specification or its complement. (I) is useful for:
CC (a) assessing whether a patient is afflicted with prostate cancer;
CC (b) monitoring the progression of prostate cancer in a patient;
CC (c) assessing the efficacy of a test compound to inhibit prostate
CC cancer in a patient;
CC (d) assessing the efficacy of a therapy for inhibiting prostate cancer
CC in a patient;
CC (e) selecting a composition for inhibiting prostate cancer in a patient;
CC (f) assessing the prostate cell carcinogenic potential of a compound;
CC (g) determining whether prostate cancer has metastasized in a patient;
CC (h) assessing the aggressiveness or indolence of prostate cancer in a
CC patient;
CC (I) is also useful as a pharmacodynamic or pharmacogenomic marker.
XX
XX Sequence 294 BP; 104 A; 26 C; 69 G; 95 T; 0 other;
SQ
Query Match 3.8%; Score 38; DB 23; Length 294;
Best Local Similarity 81.5%; Pred. No. 9.9;
Matches 44; Conservative 0; Mismatches 10; Indels 0; Gaps 0;
QY 939 AAAATGAGTGTGTTTACGTCTTGTCCACAAAAAATAAAAAAAAAA 992
DB 93 AAAAGTTTGTGTTTACCCCTTCCCTCCCAAAAAAAAAAATAAAAAA 40
RESULT 49
ABQ81848
ID ABQ81848 standard; DNA; 349980 BP.
XX
AC ABQ81848;
XX
DT 19-NOV-2002 (first entry)
XX
DE Bifidobacterium longum NCC2705 related nucleotide sequence SEQ ID:1104.
XX
KW Bifidobacterium longum NCC2705; Bifidobacterium; bacterial;
KW antidiarrheic; antibacterial; inhibitor of Salmonella; detection;
KW identification; lactic acid bacterium; diarrhoea; pathogenic bacteria;
KW rotavirus; food composition; pharmaceutical composition; gene; ds.
XX
OS Bifidobacterium longum.
OS Synthetic.
XX
PN EP1227152-A1.
XX
PD 31-JUL-2002.
XX
PF 30-JAN-2001; 2001EP-0102050.
XX
PR 30-JAN-2001; 2001EP-0102050.
XX
PA (NEST) SOC PROD NESTLE SA.
XX
DR WPI; 2002-668397/72.
XX
XX Novel polynucleotide comprising Bifidobacterium genome sequence useful
PT as a probe or primer for detecting and/or identifying Bifidobacterium
PT longum in a biological sample -
XX

PS Disclosure; SEQ ID 1104; 80pp; English.
XX
XX The present invention describes a polynucleotide (I) comprising a
CC sequence of a Bifidobacterium genome selected from the nucleotide
CC sequences given in ABQ81842 and ABQ81843, or a sequence exhibiting at
CC least 90% identity or which hybridises with the sequences given in
CC ABQ81842 and ABQ81843. Also described is a polynucleotide (II) encoding
CC a fusion protein, comprising a sequence selected from 1097 sequences
CC given in ABP65258 to ABP66354 ligated in frame to a polynucleotide
CC encoding a heterologous polypeptide. (I) has antidiarrheic and
CC antibacterial activities, and can be used as an inhibitor of Salmonella.
CC (I) (which is a probe) is useful for the detection and/or identification
CC of Bifidobacterium longum in a biological sample. A carrier containing
CC the lactic acid bacterium Bifidobacterium longum NCC2705 (NCIM 1-2618)
CC can be used for preventing and/or treating diarrhoea brought about by
CC pathogenic bacteria and/or rotavirus. The carrier is a food composition
CC selected from milk, yogurt, curd, cheese, fermented milks, milk based
CC fermented products, ice-creams, fermented cereal based products, milk
CC based powders, infant formula, pet food or a pharmaceutical composition
CC selected from tablets, liquid bacterial suspensions, dried oral
CC supplement, wet oral supplement, dry tube feeding or wet tube feeding.
CC (I) is useful in DNA arrays or chips to carry out analysis of the
CC expression of the Bifidobacterium gene. ABQ81844 to ABQ81850 represent
CC Bifidobacterium related nucleotide sequences given in the sequence
CC Listing from the present invention but not mentioned further within the
CC specification.
CC N.B. The sequence data for this patent is not represented in the printed
CC specification but is based on sequence information supplied by the
CC European Patent Office.
XX
SQ Sequence 349980 BP; 69195 A; 106952 C; 106128 G; 67705 T; 0 other;
Query Match 3.8%; Score 38; DB 24; Length 349980;
Best Local Similarity 60.8%; Pred. No. 1e+02;
Matches 62; Conservative 0; Mismatches 40; Indels 0; Gaps 0;
QY 255 ACCAAGAACATCAAGGTGGCCAGAGAGTGTGTGAAGACCCACGAGCGCGCTCTTCAAC 314
DB 222380 ACCAAGACCTTCAAGACCGACAGAGGTCCAGGCTTACTCCGAGCGCGCTCGTCATC 222439
QY 315 CTCACGCTCACACTCAAGTCCAGTCCAGCTGCTCACCTAC 356
DB 222440 CAGACCTTCACATCCCGTCTCCCTTGACCACTTCAAGAAC 222481
RESULT 50
ABV18588
ID ABV18588 standard; cDNA; 353 BP.
XX
AC ABV18588;
XX
DT 13-SEP-2002 (first entry)
XX
DE Human prostate expression marker cDNA 18579.
XX
KW Human; prostate cancer; cytostatic; carcinogen; pharmacodynamic marker;
KW pharmacogenomic marker; gene; ss.
XX
OS Homo sapiens.
XX
PN WO2001:60860-A2.
XX
PD 23-AUG-2001.
XX
PF 20-FEB-2001; 2001WO-US05171.
XX
PR 17-FEB-2000; 2000US-183319P.
PR 16-MAR-2000; 2000US-189862P.
PR 25-MAY-2000; 2000US-207454P.
PR 09-JUN-2000; 2000US-211314P.
PR 18-JUL-2000; 2000US-219007P.
PR 13-DEC-2000; 2000US-255281P.
XX

